

## ANTIBACTERIAL AGENTS

### CROSS REFERENCE

5           This application claims the benefit of U.S. Serial No. 60/405,464, filed on August 23, 2002, under 35 U.S.C. § 119(e)(i).

### FIELD OF THE INVENTION

          The present invention relates to antibacterial agents that are useful for  
10   sterilization, sanitation, antisepsis, and disinfection.

### BACKGROUND

          The inappropriate growth of a variety of bacteria has been a problem for many years. Bacteria have caused degradation of natural product materials, infection in  
15   humans and other animals, and spoilage of foods.

          Sterilization denotes the use of either physical or chemical agents to eliminate all viable bacteria from a material, while disinfection generally refers to the use of germicidal chemical agents to destroy the potential infectivity of a material. Sanitizing refers to procedures used to simply lower the bacterial content of utensils  
20   used for food. Antisepsis refers to the topical application of chemicals to a body surface to kill or inhibit pathogenic microbes. Disinfectants are widely used for skin antisepsis in preparation for surgery.

          Bacteria are the smallest organisms that contain all the machinery required for growth and self-replication. A bacterium includes a rigid cell wall surrounding the  
25   cytoplasmic membrane, which itself encloses a single naked chromosome without a nuclear membrane. The cytoplasmic membrane consists primarily of a bi-layer of lipid molecules.

          The fundamental criterion of bactericidal action is loss of the ability of the organism to propagate indefinitely, when placed in a suitable environment.  
30   Bactericidal action suggests microbe damage of various types, including the triggering of irreversible damage to the cytoplasmic cell membrane or irreversible impairment of the DNA (or viral RNA replication. Accordingly, sterilization is not identical with destruction of microbes. Additionally, it is understood that damage to nucleic acids

(DNA or RNA) is not always irreversible, as it is known that ultraviolet light-induced damage to viral nucleic acids can be repaired by enzymatic and genetic mechanisms.

## SUMMARY OF THE INVENTION

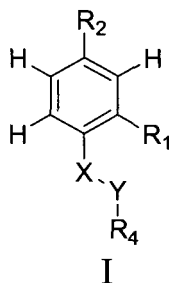
5           The invention relates to antibacterial agents that are useful for sterilization, sanitation, antiseptis, and disinfection.

          In one aspect, the invention features methods of using antibacterial agents of formula I for sterilizing, sanitizing, antiseptis, or disinfecting. The method includes applying the antibacterial agent to a location in need of sterilization, sanitation, antiseptis, and disinfection. Specifically, a method of sterilization, sanitation, antiseptis, and disinfection, includes applying antimicrobial compounds to a surface in need of sterilization, sanitation, antiseptis, and disinfection. The antimicrobial compounds are applied in a therapeutically acceptable amount, e.g., an amount sufficient to kill or hinder the growth of bacteria on the surface to be sterilized, sanitized, or disinfected.

10           antiseptis, and disinfection. Specifically, a method of sterilization, sanitation, antiseptis, and disinfection, includes applying antimicrobial compounds to a surface in need of sterilization, sanitation, antiseptis, and disinfection. The antimicrobial compounds are applied in a therapeutically acceptable amount, e.g., an amount sufficient to kill or hinder the growth of bacteria on the surface to be sterilized, sanitized, or disinfected.

15           sanitized, or disinfected.

          In general, the antibacterial agents have the formula



or a pharmaceutically acceptable salt thereof,  
wherein

20           X = NH

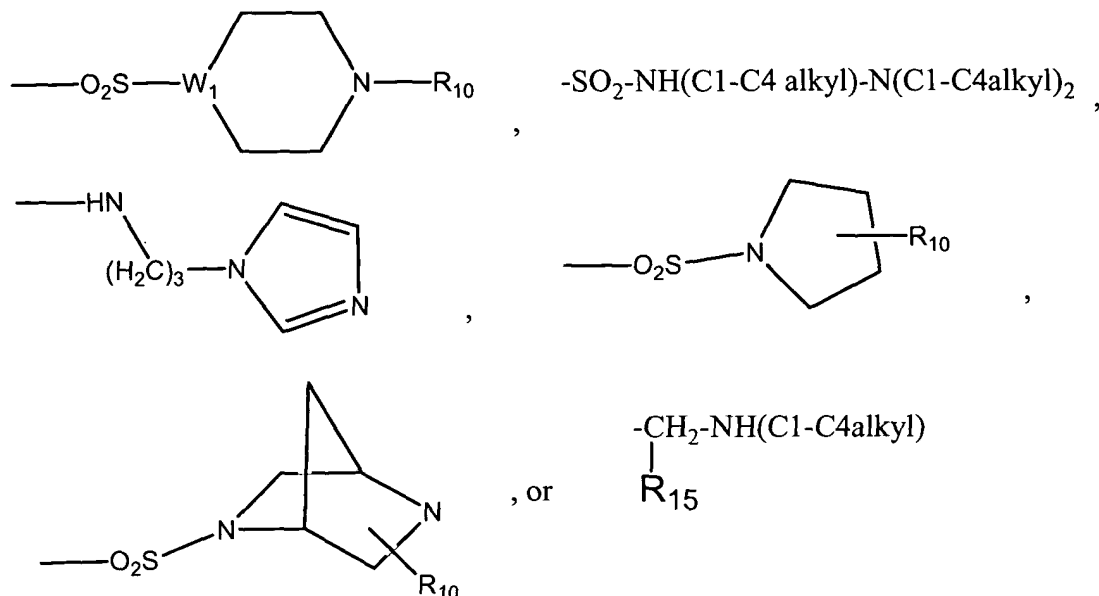
          Y = CO, CS, -C(=N-CN) or

          X and Y together form an alkene, or C<sub>3</sub>-C<sub>5</sub> cycloalkyl;

          R<sub>1</sub> is -COOH;

          R<sub>2</sub> is an electron withdrawing group;

25           R<sub>4</sub> is an optionally substituted aryl, provided that the aryl is not simultaneously substituted with a sulfonamide and a urea or thiourea, further provided that the aryl is not solely substituted at the ortho-position relative to Y, and still further provided that the aryl is not substituted with a group selected from



$W_1$  is N or CH;

$R_{10}$  is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> substituted alkyl, Het, substituted Het, aryl, or substituted aryl; and

5  $R_{15}$  is H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> substituted alkyl, Het, substituted Het, C<sub>4</sub>-C<sub>7</sub> cycloalkyl.

#### DETAILED DESCRIPTION OF THE INVENTION

The term “halo” refers to a halogen atom selected from Cl, Br, I, and F.

The term “alkyl” refers to both straight- and branched-chain moieties. Unless  
10 otherwise specifically stated alkyl moieties include between 1 and 9 carbon atoms.

The term “alkenyl” refers to both straight- and branched-chain moieties containing at least one  $-C=C-$ . Unless otherwise specifically stated alkenyl moieties include between 1 and 9 carbon atoms.

The term “alkynyl” refers to both straight- and branched-chain moieties  
15 containing at least one  $-C\equiv C-$ . Unless otherwise specifically stated alkynyl moieties include between 1 and 9 carbon atoms. between 1 and 6 carbon atoms

The term “alkoxy” refers to  $-O$ -alkyl groups.

The term “cycloalkyl” refers to a cyclic alkyl moiety. Unless otherwise specifically stated cycloalkyl moieties will include between 3 and 9 carbon atoms.

20 The term “cycloalkenyl” refers to a cyclic alkenyl moiety. Unless otherwise specifically stated cycloalkenyl moieties will include between 5 and 9 carbon atoms and at least one  $-C=C-$  group within the cyclic ring.

The term “amino” refers to  $-NH_2$ .

The term “sulfonamide” refers to a  $-S(O)_2-N(Q_{10})_2$

The term “aryl” refers to phenyl and naphthyl.

The term “het” refers to mono- or bi-cyclic ring systems containing at least one heteroatom selected from O, S, and N. Each mono-cyclic ring may be aromatic,  
 5 saturated, or partially unsaturated. A bi-cyclic ring system may include a mono-cyclic ring containing at least one heteroatom fused with an cycloalkyl or aryl group. A bi-cyclic ring system may also include a mono-cyclic ring containing at least one heteroatom fused with another het, mono-cyclic ring system.

Examples of “het” include, but are not limited to, pyridine, thiophene, furan,  
 10 pyrazoline, pyrimidine, 2-pyridyl, 3-pyridyl, 4-pyridyl, 2-pyrimidinyl, 4-pyrimidinyl, 5-pyrimidinyl, 3-pyridazinyl, 4-pyridazinyl, 3-pyrazinyl, 4-oxo-2-imidazolyl, 2-imidazolyl, 4-imidazolyl, 3-isoxazolyl, 4-isoxazolyl, 5-isoxazolyl, 3-pyrazolyl, 4-pyrazolyl, 5-pyrazolyl, 2-oxazolyl, 4-oxazolyl, 4-oxo-2-oxazolyl, 5-oxazolyl, 1,2,3-oxathiazole, 1,2,3-oxadiazole, 1,2,4-oxadiazole, 1,2,5-oxadiazole, 1,3,4-oxadiazole,  
 15 2-thiazolyl, 4-thiazolyl, 5-thiazolyl, 3-isothiazole, 4-isothiazole, 5-isothiazole, 2-furanyl, 3-furanyl, 2-thienyl, 3-thienyl, 2-pyrrolyl, 3-pyrrolyl, 3-isopyrrolyl, 4-isopyrrolyl, 5-isopyrrolyl, 1,2,3-oxathiazole-1-oxide, 1,2,4-oxadiazol-3-yl, 1,2,4-oxadiazol-5-yl, 5-oxo-1,2,4-oxadiazol-3-yl, 1,2,4-thiadiazol-3-yl, 1,2,4-thiadiazol-5-yl, 3-oxo-1,2,4-thiadiazol-5-yl, 1,3,4-thiadiazol-5-yl, 2-oxo-1,3,4-thiadiazol-5-yl,  
 20 1,2,4-triazol-3-yl, 1,2,4-triazol-5-yl, 1,2,3,4-tetrazol-5-yl, 5-oxazolyl, 3-isothiazolyl, 4-isothiazolyl, 5-isothiazolyl, 1,3,4-oxadiazole, 4-oxo-2-thiazoliny, 5-methyl-1,3,4-thiadiazol-2-yl, thiazolodione, 1,2,3,4-thiatriazole, 1,2,4-dithiazolone, phthalimide, quinolinyl, morpholinyl, benzoxazolyl, diazinyl, triazinyl, quinolinyl, quinoxalinyl, naphthyridinyl, azetidiny, pyrrolidinyl, hydantoinyl, oxathiolanyl, dioxolanyl,  
 25 imidazolidinyl, and azabicyclo[2.2.1]heptyl.

The term “heteroaryl” refers to a mono- or bicyclic het in which at least one cyclic ring is aromatic.

The term “substituted alkyl” refers to an alkyl moiety including 1-4 substituents selected from halo, het, cycloalkyl, cycloalkenyl, aryl,  $-OQ_{10}$ ,  $-SQ_{10}$ ,  $-S(O)_2Q_{10}$ ,  
 30  $-S(O)Q_{10}$ ,  $-OS(O)_2Q_{10}$ ,  $-C(=NQ_{10})Q_{10}$ ,  $-C(=N-O-Q_{10})Q_{10}$ ,  $-S(O)_2-N=S(O)(Q_{10})_2$ ,  $-S(O)_2-N=S(Q_{10})_2$ ,  $-NQ_{10}Q_{10}$ ,  $-C(O)Q_{10}$ ,  $-C(S)Q_{10}$ ,  $-C(O)OQ_{10}$ ,  $-OC(O)Q_{10}$ ,  $-C(S)NQ_{10}Q_{10}$ ,  $-N(Q_{10})C(S)NQ_{10}Q_{10}$ ,  $-C(O)NQ_{10}Q_{10}$ ,  $-C(O)C(Q_{16})_2OC(O)Q_{10}$ ,  $-CN$ ,



=O, =S, -NQ<sub>10</sub>C(O)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, and -SNQ<sub>10</sub>Q<sub>10</sub>. Each of the het, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-4 substituents independently selected from halo and Q<sub>15</sub>.

5           The term “substituted aryl” refers to an aryl moiety having 1-3 substituents selected from -OQ<sub>10</sub>, -SQ<sub>10</sub>, -S(O)<sub>2</sub>Q<sub>10</sub>, -S(O)Q<sub>10</sub>, -OS(O)<sub>2</sub>Q<sub>10</sub>, -C(=NQ<sub>10</sub>)Q<sub>10</sub>, -C(=NOQ<sub>10</sub>)Q<sub>10</sub>, -S(O)<sub>2</sub>-N=S(O)(Q<sub>10</sub>)<sub>2</sub>, -S(O)<sub>2</sub>-N=S(Q<sub>10</sub>)<sub>2</sub>, -NQ<sub>10</sub>Q<sub>10</sub>, -C(O)Q<sub>10</sub>, -C(S)Q<sub>10</sub>, -C(O)OQ<sub>10</sub>, -OC(O)Q<sub>10</sub>, -C(O)NQ<sub>10</sub>Q<sub>10</sub>, -C(O)C(Q<sub>16</sub>)<sub>2</sub>OC(O)Q<sub>10</sub>, -CN, -NQ<sub>10</sub>C(O)Q<sub>10</sub>, -N(Q<sub>10</sub>)C(S)NQ<sub>10</sub>Q<sub>10</sub>, -N(Q<sub>10</sub>)C(S)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, -SNQ<sub>10</sub>Q<sub>10</sub>, alkyl, substituted alkyl, alkenyl, alkynyl, het, halo, cycloalkyl, cycloalkenyl, and aryl. The het, cycloalkyl, cycloalkenyl, alkenyl, alkynyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>15</sub>.

15           The term “substituted het” refers to a het moiety including 1-4 substituents selected from -OQ<sub>10</sub>, -SQ<sub>10</sub>, -S(O)<sub>2</sub>Q<sub>10</sub>, -S(O)Q<sub>10</sub>, -OS(O)<sub>2</sub>Q<sub>10</sub>, -C(=NQ<sub>10</sub>)Q<sub>10</sub>, -C(=NOQ<sub>10</sub>)Q<sub>10</sub>, -S(O)<sub>2</sub>-N=S(O)(Q<sub>10</sub>)<sub>2</sub>, -S(O)<sub>2</sub>-N=S(Q<sub>10</sub>)<sub>2</sub>, -NQ<sub>10</sub>Q<sub>10</sub>, -C(O)Q<sub>10</sub>, -C(S)Q<sub>10</sub>, -C(O)OQ<sub>10</sub>, -OC(O)Q<sub>10</sub>, -C(O)NQ<sub>10</sub>Q<sub>10</sub>, -C(O)C(Q<sub>16</sub>)<sub>2</sub>OC(O)Q<sub>10</sub>, -CN, =O, =S, -NQ<sub>10</sub>C(O)Q<sub>10</sub>, -NQ<sub>10</sub>C(S)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>C(S)NQ<sub>10</sub>Q<sub>10</sub>, -S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, -SNQ<sub>10</sub>Q<sub>10</sub>, alkyl, substituted alkyl, het, halo, cycloalkyl, cycloalkenyl, and aryl. The het, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>15</sub>.

          The term “substituted alkenyl” refers to a alkenyl moiety including 1-3  
 25       substituents -OQ<sub>10</sub>, -SQ<sub>10</sub>, -S(O)<sub>2</sub>Q<sub>10</sub>, -S(O)Q<sub>10</sub>, -OS(O)<sub>2</sub>Q<sub>10</sub>, -C(=NQ<sub>10</sub>)Q<sub>10</sub>, -C(=NOQ<sub>10</sub>)Q<sub>10</sub>, -S(O)<sub>2</sub>-N=S(O)(Q<sub>10</sub>)<sub>2</sub>, -S(O)<sub>2</sub>-N=S(Q<sub>10</sub>)<sub>2</sub>, -NQ<sub>10</sub>Q<sub>10</sub>, -C(O)Q<sub>10</sub>, -C(S)Q<sub>10</sub>, -C(O)OQ<sub>10</sub>, -OC(O)Q<sub>10</sub>, -C(O)NQ<sub>10</sub>Q<sub>10</sub>, -C(S)NQ<sub>10</sub>Q<sub>10</sub>, -C(O)C(Q<sub>16</sub>)<sub>2</sub>OC(O)Q<sub>10</sub>, -CN, =O, =S, -NQ<sub>10</sub>C(S)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>C(S)NQ<sub>10</sub>Q<sub>10</sub>, -S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, -SNQ<sub>10</sub>Q<sub>10</sub>, alkyl, substituted alkyl, het, halo, cycloalkyl, cycloalkenyl, and aryl. The het, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>15</sub>.

The term “substituted alkoxy” refers to an alkoxy moiety including 1-3 substituents -OQ<sub>10</sub>, -SQ<sub>10</sub>, -S(O)<sub>2</sub>Q<sub>10</sub>, -S(O)Q<sub>10</sub>, -OS(O)<sub>2</sub>Q<sub>10</sub>, -C(=NQ<sub>10</sub>)Q<sub>10</sub>, -C(=NOQ<sub>10</sub>)Q<sub>10</sub>, -S(O)<sub>2</sub>-N=S(O)(Q<sub>10</sub>)<sub>2</sub>, -S(O)<sub>2</sub>-N=S(Q<sub>10</sub>)<sub>2</sub>, -NQ<sub>10</sub>Q<sub>10</sub>, -C(O)Q<sub>10</sub>, -C(S)Q<sub>10</sub>, -C(O)OQ<sub>10</sub>, -OC(O)Q<sub>10</sub>, -C(O)NQ<sub>10</sub>Q<sub>10</sub>, -C(S)NQ<sub>10</sub>Q<sub>10</sub>,  
 5 -C(O)C(Q<sub>16</sub>)<sub>2</sub>OC(O)Q<sub>10</sub>, -CN, =O, =S, -NQ<sub>10</sub>C(S)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>C(S)NQ<sub>10</sub>Q<sub>10</sub>, -S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, -SNQ<sub>10</sub>Q<sub>10</sub>, alkyl, substituted alkyl, het, halo, cycloalkyl, cycloalkenyl, and aryl. The het, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>15</sub>.

10 The term “substituted cycloalkenyl” refers to a cycloalkenyl moiety including 1-3 substituents -OQ<sub>10</sub>, -SQ<sub>10</sub>, -S(O)<sub>2</sub>Q<sub>10</sub>, -S(O)Q<sub>10</sub>, -OS(O)<sub>2</sub>Q<sub>10</sub>, -C(=NQ<sub>10</sub>)Q<sub>10</sub>, -C(=NOQ<sub>10</sub>)Q<sub>10</sub>, -S(O)<sub>2</sub>-N=S(O)(Q<sub>10</sub>)<sub>2</sub>, -S(O)<sub>2</sub>-N=S(Q<sub>10</sub>)<sub>2</sub>, -NQ<sub>10</sub>Q<sub>10</sub>, -C(O)Q<sub>10</sub>, -C(S)Q<sub>10</sub>, -C(O)OQ<sub>10</sub>, -OC(O)Q<sub>10</sub>, -C(O)NQ<sub>10</sub>Q<sub>10</sub>, -C(S)NQ<sub>10</sub>Q<sub>10</sub>, -C(O)C(Q<sub>16</sub>)<sub>2</sub>OC(O)Q<sub>10</sub>, -CN, =O, =S, -NQ<sub>10</sub>C(S)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)Q<sub>10</sub>,  
 15 -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>C(S)NQ<sub>10</sub>Q<sub>10</sub>, -S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, -SNQ<sub>10</sub>Q<sub>10</sub>, alkyl, substituted alkyl, het, halo, cycloalkyl, cycloalkenyl, and aryl. The het, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>15</sub>.

The term “substituted amino” refers to an amino moiety in which one or both  
 20 of the amino hydrogens are replaced with a group selected from -OQ<sub>10</sub>, -SQ<sub>10</sub>, -S(O)<sub>2</sub>Q<sub>10</sub>, -S(O)Q<sub>10</sub>, -OS(O)<sub>2</sub>Q<sub>10</sub>, -C(=NQ<sub>10</sub>)Q<sub>10</sub>, -C(=NOQ<sub>10</sub>)Q<sub>10</sub>, -S(O)<sub>2</sub>-N=S(O)(Q<sub>10</sub>)<sub>2</sub>, -S(O)<sub>2</sub>-N=S(Q<sub>10</sub>)<sub>2</sub>, -NQ<sub>10</sub>Q<sub>10</sub>, -C(O)Q<sub>10</sub>, -C(S)Q<sub>10</sub>, -C(O)OQ<sub>10</sub>, -OC(O)Q<sub>10</sub>, -C(O)NQ<sub>10</sub>Q<sub>10</sub>, -C(S)NQ<sub>10</sub>Q<sub>10</sub>, -C(O)C(Q<sub>16</sub>)<sub>2</sub>OC(O)Q<sub>10</sub>, -CN, =O, =S, -NQ<sub>10</sub>C(O)Q<sub>10</sub>, -NQ<sub>10</sub>C(S)Q<sub>10</sub>, -NQ<sub>10</sub>C(O)NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>C(S)NQ<sub>10</sub>Q<sub>10</sub>, -  
 25 S(O)<sub>2</sub>NQ<sub>10</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)<sub>2</sub>Q<sub>10</sub>, -NQ<sub>10</sub>S(O)Q<sub>10</sub>, -NQ<sub>10</sub>SQ<sub>10</sub>, -NO<sub>2</sub>, -SNQ<sub>10</sub>Q<sub>10</sub>, alkyl, substituted alkyl, het, halo, cycloalkyl, cycloalkenyl, and aryl. The het, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>15</sub>.

Each Q<sub>10</sub> is independently selected from -H, alkyl, cycloalkyl, het,  
 30 cycloalkenyl, and aryl. The het, alkyl, cycloalkyl, cycloalkenyl, and aryl being optionally substituted with 1-3 substituents selected from halo and Q<sub>13</sub>.

Each  $Q_{11}$  is independently selected from -H, halo, alkyl, aryl, cycloalkyl, and het. The alkyl, aryl, cycloalkyl, and het being optionally substituted with 1-3 substituents independently selected from halo,  $-\text{NO}_2$ ,  $-\text{CN}$ ,  $=\text{S}$ ,  $=\text{O}$ , and  $Q_{14}$ .

Each  $Q_{13}$  is independently selected from  $Q_{11}$ ,  $-\text{O}Q_{11}$ ,  $-\text{S}Q_{11}$ ,  $-\text{S}(\text{O})_2Q_{11}$ ,  
 5  $-\text{S}(\text{O})Q_{11}$ ,  $-\text{OS}(\text{O})_2Q_{11}$ ,  $-\text{C}(=\text{N}Q_{11})Q_{11}$ ,  $-\text{S}(\text{O})_2\text{N}=\text{S}(\text{O})(Q_{11})_2$ ,  $-\text{S}(\text{O})_2\text{N}=\text{S}(Q_{11})_2$ ,  
 $-\text{SC}(\text{O})Q_{11}$ ,  $-\text{N}Q_{11}Q_{11}$ ,  $-\text{C}(\text{O})Q_{11}$ ,  $-\text{C}(\text{S})Q_{11}$ ,  $-\text{C}(\text{O})\text{O}Q_{11}$ ,  $-\text{OC}(\text{O})Q_{11}$ ,  $-\text{C}(\text{O})\text{N}Q_{11}Q_{11}$ ,  
 $-\text{C}(\text{S})\text{N}Q_{11}Q_{11}$ ,  $-\text{C}(\text{O})\text{C}(Q_{16})_2\text{OC}(\text{O})Q_{10}$ ,  $-\text{CN}$ ,  $=\text{O}$ ,  $=\text{S}$ ,  $-\text{N}Q_{11}\text{C}(\text{O})Q_{11}$ ,  $-\text{N}Q_{11}\text{C}(\text{S})Q_{11}$ ,  
 $-\text{N}Q_{11}\text{C}(\text{O})\text{N}Q_{11}Q_{11}$ ,  $-\text{N}Q_{11}\text{C}(\text{S})\text{N}Q_{11}Q_{11}$ ,  $-\text{S}(\text{O})_2\text{N}Q_{11}Q_{11}$ ,  $-\text{N}Q_{11}\text{S}(\text{O})_2Q_{11}$ ,  
 $-\text{N}Q_{11}\text{S}(\text{O})Q_{11}$ ,  $-\text{N}Q_{11}\text{S}Q_{11}$ ,  $-\text{NO}_2$ , and  $-\text{SN}Q_{11}Q_{11}$ .

10 Each  $Q_{14}$  is -H or a substituent selected from alkyl, cycloalkyl, phenyl, or naphthyl, each optionally substituted with 1-4 substituents independently selected from

$-\text{F}$ ,  $-\text{Cl}$ ,  $-\text{Br}$ ,  $-\text{I}$ ,  $-\text{O}Q_{16}$ ,  $-\text{S}Q_{16}$ ,  $-\text{S}(\text{O})_2Q_{16}$ ,  $-\text{S}(\text{O})Q_{16}$ ,  $-\text{OS}(\text{O})_2Q_{16}$ ,  $-\text{N}Q_{16}Q_{16}$ ,  $-\text{C}(\text{O})Q_{16}$ ,  
 $-\text{C}(\text{S})Q_{16}$ ,  $-\text{C}(\text{O})\text{O}Q_{16}$ ,  $-\text{NO}_2$ ,  $-\text{C}(\text{O})\text{N}Q_{16}Q_{16}$ ,  $-\text{C}(\text{S})\text{N}Q_{16}Q_{16}$ ,  $-\text{CN}$ ,  $-\text{N}Q_{16}\text{C}(\text{O})Q_{16}$ ,  
 15  $-\text{N}Q_{16}\text{C}(\text{S})Q_{16}$ ,  $-\text{N}Q_{16}\text{C}(\text{O})\text{N}Q_{16}Q_{16}$ ,  $-\text{N}Q_{16}\text{C}(\text{S})\text{N}Q_{16}Q_{16}$ ,  $-\text{S}(\text{O})_2\text{N}Q_{16}Q_{16}$ , and  
 $-\text{N}Q_{16}\text{S}(\text{O})_2Q_{16}$ . The alkyl, cycloalkyl, and cycloalkenyl being further optionally substituted with  $=\text{O}$  or  $=\text{S}$ .

Each  $Q_{15}$  is alkyl, cycloalkyl, heterocycloalkyl, heteroaryl, phenyl, or naphthyl, each optionally substituted with 1-4 substituents independently selected from -F,  
 20  $-\text{Cl}$ ,  $-\text{Br}$ ,  $-\text{I}$ ,  $-\text{O}Q_{16}$ ,  $-\text{S}Q_{16}$ ,  $-\text{S}(\text{O})_2Q_{16}$ ,  $-\text{S}(\text{O})Q_{16}$ ,  $-\text{OS}(\text{O})_2Q_{16}$ ,  $-\text{C}(=\text{N}Q_{16})Q_{16}$ ,  
 $-\text{S}(\text{O})_2\text{N}=\text{S}(\text{O})(Q_{16})_2$ ,  $-\text{S}(\text{O})_2\text{N}=\text{S}(Q_{16})_2$ ,  $-\text{SC}(\text{O})Q_{16}$ ,  $-\text{N}Q_{16}Q_{16}$ ,  $-\text{C}(\text{O})Q_{16}$ ,  $-\text{C}(\text{S})Q_{16}$ ,  
 $-\text{C}(\text{O})\text{O}Q_{16}$ ,  $-\text{OC}(\text{O})Q_{16}$ ,  $-\text{C}(\text{O})\text{N}Q_{16}Q_{16}$ ,  $-\text{C}(\text{S})\text{N}Q_{16}Q_{16}$ ,  $-\text{C}(\text{O})\text{C}(Q_{16})_2\text{OC}(\text{O})Q_{16}$ , -  
 $\text{CN}$ ,  
 $-\text{N}Q_{16}\text{C}(\text{O})Q_{16}$ ,  $-\text{N}Q_{16}\text{C}(\text{S})Q_{16}$ ,  $-\text{N}Q_{16}\text{C}(\text{O})\text{N}Q_{16}Q_{16}$ ,  $-\text{N}Q_{16}\text{C}(\text{S})\text{N}Q_{16}Q_{16}$ , -  
 25  $\text{S}(\text{O})_2\text{N}Q_{16}Q_{16}$ ,  $-\text{N}Q_{16}\text{S}(\text{O})_2Q_{16}$ ,  $-\text{N}Q_{16}\text{S}(\text{O})Q_{16}$ ,  $-\text{N}Q_{16}\text{S}Q_{16}$ ,  $-\text{NO}_2$ , and  $-\text{SN}Q_{16}Q_{16}$ .  
 The alkyl, cycloalkyl, and cycloalkenyl being further optionally substituted with  $=\text{O}$  or  $=\text{S}$ .

Each  $Q_{16}$  is independently selected from -H, alkyl, and cycloalkyl. The alkyl and cycloalkyl optionally including 1-3 halos.

30 Mammal denotes human and animals.

Each  $Q_{17}$  is independently selected from -H, -OH, and alkyl optionally including 1-3 halos and -OH.

The term "electron withdrawing group" refers to the ability of a substituent to withdraw electrons relative to that of hydrogen if the hydrogen atom occupied the same position on the molecule. The term "electron withdrawing group" is well understood by one skilled in the art and is discussed in Advanced Organic Chemistry by J. March, John Wiley & Sons, New York, New York, (1985) and the discussion therein is incorporated herein by reference. Electron withdrawing groups include, but are not limited to, groups such as halo, nitro, carboxy, cyano, aryl optionally substituted, aromatic het (excluding pyridine) optionally substituted,  $-\text{OC}(\text{Z}_n)_3$ ,  $-\text{C}(\text{Z}_n)_3$ ,  $-\text{C}(\text{Z}_n)_2-\text{O}-\text{C}(\text{Z}_m)_3$ ,  $-(\text{CO})-\text{Q}_{17}$ ,  $-\text{SO}_2-\text{C}(\text{Z}_n)_3$ ,  $-\text{SO}_2\text{-aryl}$ ,  $-\text{C}(\text{NQ}_{17})\text{Q}_{17}$ ,  $-\text{CH}=\text{C}(\text{Q}_{17})_2$ ,  $-\text{C}\equiv\text{C}-\text{Q}_{17}$ , in which each  $\text{Z}_n$  and  $\text{Z}_m$  is independently H, halo,  $-\text{CN}$ ,  $-\text{NO}_2$ ,  $-\text{OH}$ , or  $\text{C}_{1-4}\text{alkyl}$  optionally substituted with 1-3 halo,  $-\text{OH}$ ,  $\text{NO}_2$ , and provided that at least one of  $\text{Z}_n$  is halo,  $-\text{CN}$ , or  $\text{NO}_2$ , and further provided that  $\text{Q}_{17}$  is not  $-\text{OH}$  when the the electron withdrawing group is  $-(\text{CO})-\text{Q}_{17}$ .

It is to be understood that the present invention encompasses any racemic, optically-active, polymorphic, tautomeric, or stereoisomeric form, or mixture thereof, of a compound of the invention, which possesses the useful properties described herein.

In cases where compounds are sufficiently basic or acidic to form stable nontoxic acid or base salts, use of the compounds as pharmaceutically acceptable salts may be appropriate. Examples of pharmaceutically acceptable salts which are within the scope of the present invention include organic acid addition salts formed with acids which form a physiological acceptable anion and inorganic salts. Examples of pharmaceutically acceptable salts include, but are not limited to, the following acids acetic, aspartic, benzenesulfonic, benzoic, bicarbonic, bisulfuric, bitartaric, butyric, calcium edetate, camsylic, carbonic, chlorobenzoic, citric, edetic, edisylic, estolic, esyl, esylic, formic, fumaric, gluceptic, gluconic, glutamic, glycolylarsanilic, hexamic, hexylresorcinoic, hydrabamic, hydrobromic, hydrochloric, hydroiodic, hydroxynaphthoic, isethionic, lactic, lactobionic, maleic, malic, malonic, mandelic, methanesulfonic, methylnitric, methylsulfuric, mucic, muconic, napsylic, nitric, oxalic, p-nitromethanesulfonic, pamoic, pantothenic, phosphoric, monohydrogen phosphoric, dihydrogen phosphoric, phthalic, polygalactouronic, propionic, salicylic, stearic, succinic, sulfamic, sulfanilic, sulfonic, sulfuric, tannic, tartaric, teoclic toluenesulfonic, primary, secondary, and tertiary amines, substituted amines including

naturally occurring substituted amines, cyclic amines, such as arginine, betaine, caffeine, choline, N, N-dibenzylethylenediamine, diethylamine, 2-diethylaminoethanol, 2-dimethylamino-ethanol, ethanolamine, ethylenediamine, N-ethylmorpholine, N-ethylpiperidine, glucamine, glucosamine, histidine, hydrabamine, isopropylamine, lysine, methylglucamine, morpholine, piperazine, piperidine, polyamine resins, procaine, purines, theobromine, triethylamine, trimethylamine, tripropylamine, and the like.

Pharmaceutically acceptable salts may be obtained using standard procedures well known in the art, for example by reacting a sufficiently basic compound such as an amine with a suitable acid affording a physiologically acceptable anion. Alkali metal (for example, sodium, potassium or lithium) or alkaline earth metal (for example calcium) salts of carboxylic acids can also be made.

The antibacterial agents of this invention have useful activity against a variety of organisms. The in vitro activity of compounds of this invention can be assessed by standard testing procedures such as the determination of minimum inhibitory concentration (MIC) by agar dilution as described in "Approved Standard. Methods for Dilution Antimicrobial Susceptibility Tests for Bacteria That Grow Aerobically", 3rd. ed., published 1993 by the National Committee for Clinical Laboratory Standards, Villanova, Pennsylvania, USA.

The antibacterial agents described herein are useful for sterilization, sanitation, antisepsis, and disinfection. The antibacterial agents can be applied to a location in need of sterilization, sanitation, antisepsis, or disinfection, by methods known to those skilled in the art. For instance, the antibacterial agents may be incorporated into a cleaning solution that is applied, such as by spraying or pouring, to an item in need of sterilization, sanitation, antisepsis, or disinfection. The antibacterial agents may be used alone or in combination, e.g., agents disclosed herein with one another or agent(s) disclosed herein with other antibacterial agents. The antibacterial agents may be applied in varying concentrations depending upon the bacterial susceptibility to antibacterial agent(s) being applied and the desired level of sterilization, sanitation, antisepsis, or disinfection.

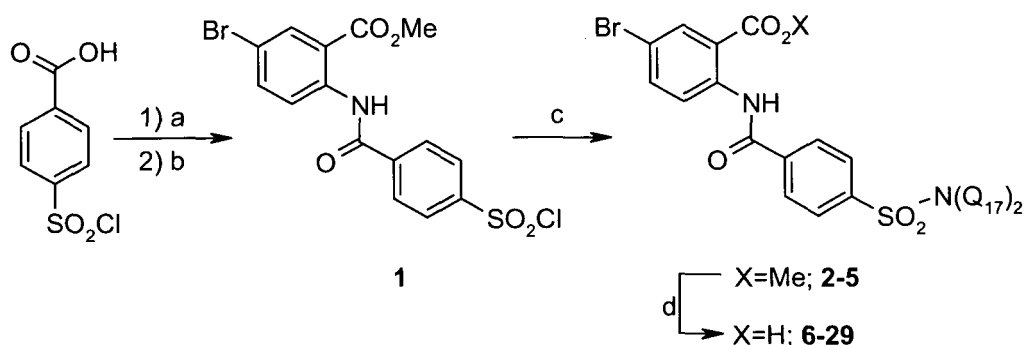
The antibacterial compounds of this invention may be synthesized by various methods known to those skilled in the art. Non-limiting examples of synthetic schemes for producing the antibacterial agents are described below.

## EXAMPLES

Without further elaboration, it is believed that one skilled in the art can, using the preceding description, practice the present invention to its fullest extent. The following detailed examples describe how to prepare the various compounds and/or perform the various processes of the invention and are to be construed as merely illustrative, and not limitations of the preceding disclosure in any way whatsoever. Those skilled in the art will promptly recognize appropriate variations from the procedures both as to reactants and as to reaction conditions and techniques.

### Example 1: Sulfonyl Derivatives

#### Scheme 1.1



a) oxalyl chloride; b) Methyl-2-amino-5-bromobenzoate; c) HN(Q<sub>17</sub>)<sub>2</sub>; d) KOH

#### Methyl 5-bromo-2-[[4-(chlorosulfonyl)benzoyl]amino]benzoate

Methyl 5-bromo-2-[[4-(chlorosulfonyl)benzoyl]amino]benzoate (**1**) was prepared as a common intermediate for the formation of sulfonamides by the procedure below: 4-(chlorosulfonyl)benzoic acid (18.37 g, 8.33 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (140 mL) and 4 drops of DMF. The solution was cooled to 0° C and oxalyl chloride (1.8 mL, 20.6 mmol) was added and stirred for 1 hour, removed from ice bath, and stirred overnight. The clear solution was concentrated *in vacuo*, redissolved in CH<sub>2</sub>Cl<sub>2</sub>, and concentrated *in vacuo*. The resulting product was dissolved in toluene (140 mL) and refluxed for 30 minutes to remove any HCl gas. After cooling to room temperature, methyl-2-amino-5-bromobenzoate (15.96 g, 69.4 mmol) was added, and the

suspension was refluxed overnight. The suspension was cooled to 0° C and filtered, washing with toluene and quickly with ethyl acetate. The solid was dried in a vacuum oven overnight to obtain sulfonyl chloride 1 (19.8 g, 66%). <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 12.19, 8.82, 8.27-8.19, 7.73, 4.00; IR 1700, 1683, 1604, 1585, 1524 (s), cm<sup>-1</sup>; MS (ESI-) for C<sub>15</sub>H<sub>11</sub>BrClNO<sub>5</sub>S *m/z* 429.8 (M-H)<sup>-</sup>.

#### General Method A (sulfonamide preparation with anilines, primary, and secondary amines)

##### 10 Methyl 5-bromo-2-({4-[(diethylamino)sulfonyl]benzoyl}amino)benzoate.

To a solution of the sulfonyl chloride 1, (694.1 mg, 1.61 mmol, 1.0 eq) in toluene (4.0 mL) was added diethyl amine (500 μL, 4.83 mmol, 3.0 eq). The suspension was shaken at 50° C for overnight. The product was extracted with EtOAc, washed with 1 N HCl and water, and concentrated *in vacuo*. The compound was dried in a vacuum oven at 50° C overnight to obtain 624.4 mg (83%). <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 11.48, 8.31, 8.11, 8.05, 7.99, 7.87, 3.86, 3.20, 1.04; IR 1700, 1676 (s), 1600, 1519 (s), 1338, 1330, 1306 (s), cm<sup>-1</sup>. Anal. Calcd for C<sub>19</sub>H<sub>21</sub>BrN<sub>2</sub>O<sub>5</sub>S: C, 48.62; H, 4.51; N, 5.97; Br, 17.02; S, 6.83. Found: C, 48.76; H, 4.53; N, 5.89; Br, 16.98; S, 6.73.

##### 20 General Method B (hydrolysis of the methyl ester)

##### 5-bromo-2-({4-[(diethylamino)sulfonyl]benzoyl}amino)benzoate, 8.

Methyl 5-bromo-2-({4-[(diethylamino)sulfonyl]benzoyl}amino)benzoate (329.6 mg, 0.704 mmol) was dissolved in 2 mL of dioxane and 0.2 mL of water. KOH (1 pellet, ~80 mg) was added to the mixture as it was heated at 50° C for 3 hours. The reaction was cooled, extracted with EtOAc, washed with 1 N HCl and brine, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated *in vacuo*, and dried in a vacuum oven at 50° C overnight to yield 313.8 mg (98%). <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.05, 8.55, 8.11, 8.09, 8.00, 7.86, 3.19, 1.04; IR 1703, 1661, 1202, 1185, cm<sup>-1</sup>. MS (FAB) *m/z* (rel. intensity) 455 (MH<sup>+</sup>, 45), 457 (37), 455 (45), 240 (99). HRMS (FAB) calcd for

C<sub>18</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>5</sub>S + H<sub>1</sub> 455.0276, found 455.0260. Anal. Calcd for C<sub>18</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>5</sub>S: C, 47.48; H, 4.21; N, 6.15; Br, 17.55; S, 7.04. Found: C, 47.31; H, 4.25; N, 6.12.

**5-bromo-2-({4-[(dimethylamino)sulfonyl]benzoyl}amino)benzoic acid 6**, was prepared by method B from its methyl ester, i.e., Methyl 5-bromo-2-({4-

5 [(dimethylamino)sulfonyl]benzoyl}amino)benzoate, in a 47% yield. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 8.89, 8.31, 8.18, 7.96, 7.78, 2.78; IR 3135, 1700, 1350 (s), 1191 (s), cm<sup>-1</sup>. MS (ESI-) for C<sub>16</sub>H<sub>15</sub>BrNO<sub>5</sub>S *m/z* 426.9 (M-H, Br isotope)<sup>-</sup>. Anal. Calcd for C<sub>16</sub>H<sub>15</sub>BrN<sub>2</sub>O<sub>5</sub>S: C, 44.98; H, 3.54; N, 6.56; Br, 18.70; S, 7.50. Found: C, 44.82; H, 3.55; N, 6.46; Br, 18.43; S, 7.36.

10 **5-bromo-2-({4-[(1H-indol-5-ylamino)sulfonyl]benzoyl}amino)benzoate 7**, was prepared by general method B from PNU-263551 in a 52% yield. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.05 (s, 1 H), 11.05 (s, 1 H), 10.00 (s, 1 H), 8.52 (d, *J* = 9 Hz, 1 H), 8.10 (d, *J* = 2 Hz, 1 H), 8.02 (d, *J* = 8 Hz, 2 H), 7.85 (m, 3 H), 7.30 (t, *J* = 1 Hz, 1 H), 7.25440 (s, 1 H), 7.24 (d, *J* = 9 Hz, 1 H), 6.82 (dd, *J* = 9, 1 Hz, 1 H), 6.34 (s, 1 H);  
15 IR 1687, 1664, 1607, 1524, 1338, 1314, 1300, 1189, 1170 (s), 825, 801, 756, 743, 681, 616 (s), cm<sup>-1</sup>. MS (FAB) *m/z* (rel. intensity) 514 (MH<sup>+</sup>, 55), 516 (59), 515 (67), 514 (55), 132 (99), 131 (97). HRMS (FAB) calcd for C<sub>22</sub>H<sub>16</sub>BrN<sub>3</sub>O<sub>5</sub>S + H<sub>1</sub> 514.0073, found 514.0066. HPLC [1] shows one main peak at 16.3 min (95%). Anal. Calcd for C<sub>22</sub>H<sub>16</sub>BrN<sub>3</sub>O<sub>5</sub>S: C, 51.37; H, 3.13; N, 8.17; Br, 15.53; S, 6.23. Found: C,  
20 51.16; H, 3.23; N, 8.01.

**5-bromo-2-[(4-{[(3-furylmethyl)amino]sulfonyl}benzoyl)amino]benzoate 9**, was prepared by method B from PNU-276173 in a 48% yield. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.60 (d, *J* = 9 Hz, 1 H), 8.41 (t, *J* = 6 Hz, 1 H), 8.14 (d, *J* = 2 Hz, 1 H), 8.07 (d, *J* = 8 Hz, 2 H), 7.93 (d, *J* = 8 Hz, 2 H), 7.87 (dd, *J* = 9, 2 Hz, 1 H), 7.46 (s, 1  
25 H), 6.28 (s, 1 H), 6.18 (s, 1 H), 4.08 (d, *J* = 6 Hz, 2 H); IR 3252, 1702, 1172 (s), 1165 (s), cm<sup>-1</sup>. MS (FAB) *m/z* (rel. intensity) 479 (MH<sup>+</sup>, 13), 481 (14), 479 (13), 135 (99), 73 (64). HRMS (FAB) calcd for C<sub>19</sub>H<sub>15</sub>BrN<sub>2</sub>O<sub>6</sub>S + H<sub>1</sub> 478.9913, found 478.9922. Anal. Calcd for C<sub>19</sub>H<sub>15</sub>BrN<sub>2</sub>O<sub>6</sub>S: C, 47.61; H, 3.15; N, 5.84; Br, 16.67; S, 6.69. Found: C, 47.55; H, 3.22; N, 5.69; Br, 16.26; S, 6.60.

30 **5-bromo-2-[(4-{[4-(ethoxycarbonyl)-1-piperazinyl]sulfonyl}benzoyl)amino]benzoic acid 10** was prepared by method A followed by B with a 26% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>)



$\delta$  8.60 (d,  $J$  = 9 Hz, 1 H), 8.18 (d,  $J$  = 8 Hz, 2 H), 8.13 (d,  $J$  = 2 Hz, 1 H), 7.94 (d,  $J$  = 8 Hz, 2 H), 7.79 (dd,  $J$  = 9, 2 Hz, 1 H), 3.97 (q,  $J$  = 7 Hz, 2 H), 3.45 (br. s, 4 H), 2.95 (br. s, 4 H), 1.12 (t,  $J$  = 7 Hz, 3 H); IR 1692 (s), 1675 (s), 1584, 1518 (s), 1287, 1276, 1250,  $\text{cm}^{-1}$ . MS (FAB)  $m/z$  (rel. intensity) 540 ( $\text{MH}^+$ , 46), 542 (44), 540 (46), 159 (95), 157 (99). HRMS (FAB) calcd for  $\text{C}_{21}\text{H}_{22}\text{BrN}_3\text{O}_7\text{S} + \text{H}_1$  540.0440, found 540.0428. HPLC [1] shows one major peak at 16.2 min (97%). Anal. Calcd for  $\text{C}_{21}\text{H}_{22}\text{BrN}_3\text{O}_7\text{S}$ : C, 46.67; H, 4.10; N, 7.78; Br, 14.79; S, 5.93. Found: C, 46.34; H, 4.19; N, 7.63; Br, 14.18; S, 5.79.

**5-bromo-2-([4-([methyl[2-(2-pyridinyl)ethyl]amino)sulfonyl]benzoyl]amino) benzoic acid 11** was prepared by method A followed by B with a 57% yield over both steps. The methyl ester was not fully characterized.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.19 (s, 1 H), 8.58 (d,  $J$  = 9 Hz, 1 H), 8.52 (d,  $J$  = 4 Hz, 1 H), 8.13 (d,  $J$  = 3 Hz, 1 H), 8.12 (d,  $J$  = 6 Hz, 2 H), 7.96 (d,  $J$  = 8 Hz, 2 H), 7.87 (dd,  $J$  = 9, 2 Hz, 1 H), 7.78 (td,  $J$  = 8, 2 Hz, 1 H), 7.35 (d,  $J$  = 8 Hz, 1 H), 7.30 (td,  $J$  = 6, 2 Hz, 1 H), 3.42 (t,  $J$  = 7 Hz, 2 H), 2.99 (t,  $J$  = 8 Hz, 2 H), 2.77 (s, 3 H); IR 1692 (s), 1518 (s), 1340 (s), 1297 (s), 1162 (s), 763 (s), 755 (s), 747 (s)  $\text{cm}^{-1}$ . MS (ES-) for  $\text{C}_{22}\text{H}_{20}\text{BrN}_3\text{O}_5\text{S}$   $m/z$  518.0 ( $\text{M}-\text{H}^+$ , Br isotope); HRMS (FAB) calcd for  $\text{C}_{22}\text{H}_{20}\text{BrN}_3\text{O}_5\text{S} + \text{H}_1$  518.0386, found 518.0388. HPLC [1] shows one major peak (13.58 min, 99%).

**2-([4-([benzylamino)sulfonyl]benzoyl]amino)-5-bromobenzoic acid 12** was prepared by method A followed by B with a 17% yield over both steps. The methyl ester was not fully characterized.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.09 (s, 1 H), 8.60 (d,  $J$  = 9 Hz, 1 H), 8.39 (t,  $J$  = 6 Hz, 1 H), 8.14 (d,  $J$  = 2 Hz, 1 H), 8.08 (d,  $J$  = 8 Hz, 2 H), 7.97 (d,  $J$  = 8 Hz, 2 H), 7.88 (dd,  $J$  = 9, 2 Hz, 1 H), 7.30-7.20 (m, 5 H), 4.05 (d,  $J$  = 6 Hz, 2 H); HRMS (FAB) calcd for  $\text{C}_{21}\text{H}_{17}\text{BrN}_2\text{O}_5\text{S} + \text{H}_1$  489.0120, found 489.0129; HPLC [1] shows one major peak (20.60 min, 99%).

**5-bromo-2-([4-([2-hydroxy-1-methylethyl]amino)sulfonyl]benzoyl]amino) benzoic acid 14** was prepared by method A followed by B with a 35% yield over both steps. The methyl ester was not fully characterized.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ )  $\delta$  8.56 (d,  $J$  = 9 Hz, 1 H), 8.11 (d,  $J$  = 2 Hz, 1 H), 8.09 (d,  $J$  = 8 Hz, 2 H), 8.00 (d,  $J$  = 8 Hz, 2 H), 7.86 (dd,  $J$  = 9, 2 Hz, 1 H), 7.76 (d,  $J$  = 7 Hz, 1 H), 3.26 (m, 2 H), 3.12 (m, 1 H), 0.89 (d,  $J$  = 6 Hz, 3 H); MS (ES-) for  $\text{C}_{17}\text{H}_{17}\text{BrN}_2\text{O}_6\text{S}$   $m/z$  454.9 ( $\text{M}-\text{H}^+$ ); HPLC [1] shows one major peak (14.08 min, 96%).

**5-bromo-2-({4-[(4-carboxyanilino)sulfonyl]benzoyl}amino)benzoic acid 15** was prepared from method A followed by method B in a 10% yield. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.45 (br. s, 1 H), 11.15 (s, 1 H), 8.52 (d, *J* = 9 Hz, 1 H), 8.08 (d, *J* = 8 Hz, 3 H), 8.03 (d, *J* = 9 Hz, 2 H), 7.81 (d, *J* = 9 Hz, 3 H), 7.26 (d, *J* = 9 Hz, 2 H); HPLC [1] shows one major peak (15.15 min, 90%).

**5-bromo-2-{[4-(3,4-dihydro-1(2H)-quinolinylsulfonyl)benzoyl]amino}benzoic acid 16** was prepared by method A followed by method B in a 48% yield. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.05 (s, 1 H), 8.52 (d, *J* = 9 Hz, 1 H), 8.11 (d, *J* = 3 Hz, 1 H), 8.05 (d, *J* = 9 Hz, 2 H), 7.86 (dd, *J* = 9, 2 Hz, 1 H), 7.82 (d, *J* = 8 Hz, 2 H), 7.61 (d, *J* = 8 Hz, 1 H), 7.25-7.05 (m, 3 H), 3.83 (t, *J* = 6 Hz, 2 H), 2.45 (t, *J* = 7 Hz, 2 H), 1.63 (quintet, *J* = 6 Hz, 2 H); IR 1667, 1601, 1584, cm<sup>-1</sup>. HRMS (FAB) calcd for C<sub>23</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>5</sub>S + H<sub>1</sub> 515.0276, found 515.0264. Anal. Calcd for C<sub>23</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>5</sub>S: C, 53.60; H, 3.72; N, 5.43. Found: C, 53.52; H, 3.96; N, 5.57.

**5-bromo-2-{[4-({2-(3,5-dimethoxyphenyl)ethyl}amino)sulfonyl]benzoyl}amino}benzoic acid 17** was prepared by method A followed by B with a 56% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.60 (d, *J* = 9 Hz, 1 H), 8.13 (d, *J* = 3 Hz, 1 H), 8.09 (d, *J* = 8 Hz, 2 H), 7.95 (d, *J* = 9 Hz, 2 H), 7.87 (dd, *J* = 9, 2 Hz, 1 H), 6.79 (d, *J* = 8 Hz, 1 H), 6.73 (d, *J* = 2 Hz, 1 H), 6.64 (dd, *J* = 8, 2 Hz, 1 H), 3.70 (s, 3 H), 3.68 (s, 3 H), 3.02 (q, *J* = 6 Hz, 2 H), 2.61 (t, *J* = 7 Hz, 2 H); MS (FAB) *m/z* (rel. intensity) 563 (MH<sup>+</sup>, 86), 565 (86), 564 (82), 563 (86), 562 (56), 348 (77), 199 (46), 165 (56), 164 (32), 152 (49), 151 (99). HRMS (EI) calcd for C<sub>24</sub>H<sub>23</sub>BrN<sub>2</sub>O<sub>7</sub>S 562.0410, found 562.0438. HPLC [1] shows one major peak (16.16 min, 97%).

**5-bromo-2-[4-({(3S)-3-hydroxypyrrolidinyl}sulfonyl)benzoyl]amino}benzoic acid 13** was prepared by method A followed by B in a 15% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.62 (d, *J* = 9 Hz, 1 H), 8.18 (d, *J* = 8 Hz, 2 H), 8.11 (d, *J* = 3 Hz, 1 H), 7.92 (d, *J* = 11 Hz, 2 H), 7.78 (dd, *J* = 9, 2 Hz, 1 H), 5.16 (m, 1 H), 3.50-3.20 (m, 4 H), 2.10-1.90 (m, 2 H); HPLC [1] shows one major peak (18.94 min, 97%).

**5-bromo-2-({4-[(ethylanilino)sulfonyl]benzoyl}amino)benzoic acid 19** was

prepared by method A followed by B with a 75% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, CD<sub>3</sub>OD) δ 8.75 (d, *J* = 9 Hz, 1 H), 8.24 (d, *J* = 2 Hz, 1 H), 8.11 (d, *J* = 8 Hz, 2 H), 7.76 (dd, *J* = 9, 2 Hz, 1 H), 7.74 (d, *J* = 8 Hz, 2 H), 7.34 (m, 3 H), 7.06 (m, 2 H), 3.69 (q, *J* = 7 Hz, 2 H), 1.07 (t, *J* = 7 Hz, 3 H); MS (ES<sup>-</sup>) for C<sub>22</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>5</sub>S *m/z* 502.8 (M-H<sup>+</sup>; Br isotope); HRMS (FAB) calcd for C<sub>22</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>5</sub>S +H<sub>1</sub> 503.0276, found 503.0265. HPLC [1] shows one major peak (18.60 min, 99%).

**5-bromo-2-({4-[(3,5-dimethoxyanilino)sulfonyl]benzoyl}amino)benzoic acid 20**

was prepared by method A followed by B with a 69% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, CD<sub>3</sub>OD) δ 8.73 (d, *J* = 9 Hz, 1 H), 8.24 (d, *J* = 2 Hz, 1 H), 8.09 (d, *J* = 9 Hz, 2 H), 7.96 (d, *J* = 9 Hz, 2 H), 7.74 (dd, *J* = 9, 2 Hz, 1 H), 6.32 (s, 1 H), 6.31 (s, 1 H), 6.20 (s, 1 H), 3.70 (s, 6 H); MS (ES<sup>-</sup>) for C<sub>22</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>7</sub>S *m/z* 532.8 (M-H<sup>+</sup>); HPLC [1] shows one major peak (17.06 min, 96%).

**5-bromo-2-[(4-[(2-hydroxy-2-phenylethyl)(methyl)amino]sulfonyl]**

**benzoyl]amino] benzoic acid 21** was prepared by method A followed by B with a 15% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, CD<sub>3</sub>OD) δ 12.10 (s, 1 H), 8.57 (d, *J* = 9 Hz, 1 H), 8.12 (d, *J* = 2 Hz, 1 H), 8.11 (d, *J* = 9 Hz, 2 H), 7.95 (d, *J* = 8 Hz, 2 H), 7.87 (dd, *J* = 9, 3 Hz, 1 H), 7.35-7.27 (m, 5 H), 4.76 (t, *J* = 7 Hz, 1 H), 3.22-3.13 (m, 2 H), 2.77 (s, 3 H); MS (FAB) *m/z* (rel. intensity) 533 (MH<sup>+</sup>, 61), 535 (64), 533 (61), 517 (99), 516 (24), 515 (90), 318 (46), 152 (27), 134 (25), 132 (33), 44 (44). HRMS (FAB) calcd for C<sub>23</sub>H<sub>21</sub>BrN<sub>2</sub>O<sub>6</sub>S +H<sub>1</sub> 533.0382, found 533.0386. HPLC [1] shows one major peak (17.06, 97%).

**5-bromo-2-{{4-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl}amino}benzoic acid 22**

was prepared by method A followed by B in a 55% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.00 (s, 1 H), 8.51 (d, *J* = 9 Hz, 1 H), 8.10-8.01 (m, 5 H), 7.84 (dd, *J* = 9, 3 Hz, 1 H), 7.50 (d, *J* = 8 Hz, 1 H), 7.22 (t, *J* = 8 Hz, 1 H), 7.17 (d, *J* = 8 Hz, 1 H), 7.00 (t, *J* = 7 Hz, 1 H), 3.98 (t, *J* = 8 Hz, 2 H), 2.93 (t, *J* = 8 Hz, 2 H); IR 1687, 1667, 1601, 1525 (s), 1365 (s), 1245 (s), 1172 (s), cm<sup>-1</sup>. MS (FAB) *m/z* (rel. intensity) 501 (MH<sup>+</sup>, 36), 503 (41), 502 (43), 501 (36), 500 (31), 286 (35), 118 (99). HRMS (FAB) calcd for

C<sub>22</sub>H<sub>17</sub>BrN<sub>2</sub>O<sub>5</sub>S + H<sub>2</sub>O 501.0120, found 501.0118. Anal. Calcd for C<sub>22</sub>H<sub>17</sub>BrN<sub>2</sub>O<sub>5</sub>S: C, 52.71; H, 3.42; N, 5.59; Br, 15.94; S, 6.39. Found: C, 52.65; H, 3.47; N, 5.58; Br, 15.88; S, 6.24.

**5-bromo-2-({4-[(5-methoxy-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)**

5 **benzoic acid 23** was prepared by method A followed by B in a 17% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.05 (s, 1 H), 8.51 (d, *J* = 9 Hz, 1 H), 8.10 (d, *J* = 2 Hz, 1 H), 8.05 (d, *J* = 8 Hz, 2 H), 7.95 (d, *J* = 9 Hz, 2 H), 7.86 (dd, *J* = 9, 2 Hz, 1 H), 7.42 (d, *J* = 9 Hz, 1 H), 6.78 (d, *J* = 8 Hz, 1 H), 6.77 (s, 1 H), 3.96 (t, *J* = 8 Hz, 2 H), 3.68 (s, 3 H), 2.80 (t, *J* = 8 Hz, 2 H); IR 1702, 1606, 1518, 1489 (s), 1358, 1199 (s), 1168 (s), cm<sup>-1</sup>. MS (FAB) *m/z* (rel. intensity) 531 (MH<sup>+</sup>, 29), 533 (30), 531 (29), 530 (38), 148 (99). HRMS (EI) calcd for C<sub>23</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>6</sub>S 530.0148, found 530.0156. Anal. Calcd for C<sub>23</sub>H<sub>19</sub>BrN<sub>2</sub>O<sub>6</sub>S: C, 51.99; H, 3.60; N, 5.27; Br, 15.04; S, 6.03. Found: C, 52.08; H, 3.61; N, 5.29.

15 **5-bromo-2-({4-[(5-fluoro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)**

**benzoic acid 24** was prepared by method A followed by B with a 41% yield over both steps. The methyl ester was not fully characterized. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 12.05 (s, 1 H), 8.51 (d, *J* = 9 Hz, 1 H), 8.10 (d, *J* = 2 Hz, 1 H), 8.07 (d, *J* = 9 Hz, 2 H), 7.99 (d, *J* = 9 Hz, 2 H), 7.85 (dd, *J* = 9, 2 Hz, 1 H), 7.49 (dd, *J* = 10, 5 Hz, 1 H), 7.07-7.02 (m, 2 H), 4.01 (t, *J* = 8 Hz, 2 H), 2.89 (t, *J* = 8 Hz, 2 H); MS (ES<sup>-</sup>) for C<sub>22</sub>H<sub>16</sub>BrN<sub>2</sub>O<sub>5</sub>S *m/z* 518.9 (M-H<sup>+</sup>, Br isotope); HPLC [2] shows one major peak (6.35 min, 96%).

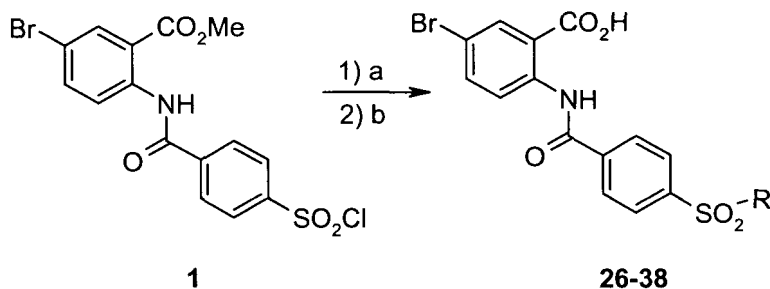
**2-{{4-[(1H-benzimidazol-1-yl)sulfonyl]benzoyl}amino}-5-bromobenzoic acid 26** was

prepared from method A followed by hydrolysis of the methyl ester by the hydrolysis  
25 procedure in method C below. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 11.98 (s, 1 H), 8.91 (s, 1 H), 8.47 (d, *J* = 9 Hz, 1 H), 8.41 (d, *J* = 9 Hz, 2 H), 8.13 (d, *J* = 9 Hz, 2 H), 8.09 (d, *J* = 2 Hz, 1 H), 7.93 (d, *J* = 7 Hz, 1 H), 7.85 (dd, *J* = 9, 3 Hz, 1 H), 7.78 (d, *J* = 7 Hz, 1 H), 7.47 (t, *J* = 6 Hz, 1 H), 7.40 (t, *J* = 6 Hz, 1 H); IR 1686, 1607, 1522, 1391, 1296, 1262, 1190, cm<sup>-1</sup>. MS (ESI<sup>-</sup>) for C<sub>21</sub>H<sub>14</sub>BrN<sub>3</sub>O<sub>5</sub>S *m/z* 497.7 (M-H)<sup>-</sup>. HPLC [2]  
30 shows one major peak at 6.01 min (96%). Anal. Calcd for C<sub>21</sub>H<sub>14</sub>BrN<sub>3</sub>O<sub>5</sub>S: C, 50.21; H, 3.21; N, 8.36; Br, 15.91; S, 6.38. Found: C, 50.06; H, 2.85; N, 7.93; Br, 15.34; S, 6.22.

### General Method C (sulfonamide preparation with indoles and pyrrole):

Reaction of sulfonyl chloride intermediate **1** with indole derivatives requires modified conditions. Deprotonation of the indole nitrogen with sodium hydride in THF and reaction with the sulfonyl chloride **1** provided the desired intermediate methyl esters. Two equivalents of the indole anion were required because of competitive deprotonation of the amide in **1**. Attempted hydrolysis of such methyl esters with aqueous KOH results in hydrolysis of the newly formed sulfonamide. Therefore, dealkylative deesterification conditions were utilized (Scheme 1.2).

#### Scheme 1.2



a) R<sup>\*</sup>, NaH, THF; b) MeI, NaCN

\* R = indoles, pyrrole, indazole, and benzoxazolinone

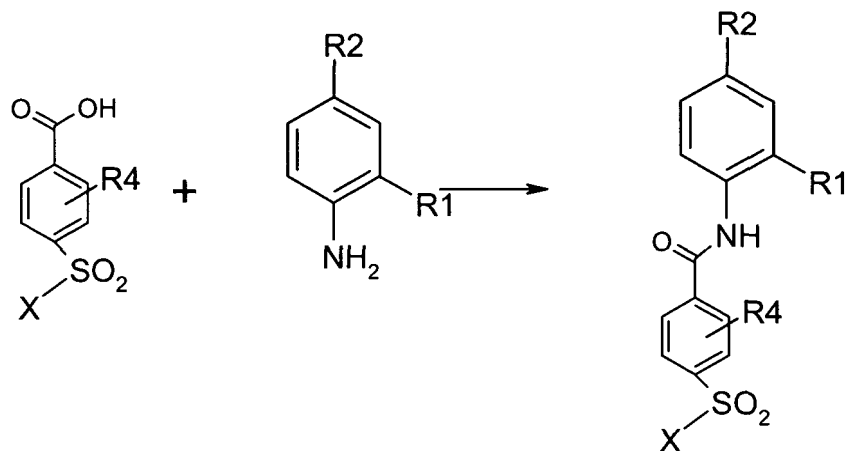
**5-bromo-2-([4-[(5-fluoro-1H-indol-1-yl)sulfonyl]benzoyl]amino)benzoic acid 26** was prepared by the following procedure: 5-fluoroindole (497.1 mg, 3.68 mmol, 2.2 eq) was dissolved in anhydrous THF (8 mL) and cooled to 0<sup>0</sup> C. NaH (60% dispersion in mineral oil, 150 mg, 3.75 mmol, 2.2 eq) was added and the cloudy mixture was stirred for 1 hr. at 0-25<sup>0</sup> C. The suspension was then cooled to 0<sup>0</sup> C and Methyl 5-bromo-2-([4-(chlorosulfonyl)benzoyl]amino)benzoate (722.0 mg, 1.68 mmol, 1.0 eq) was added neat and stirred overnight at room temperature. After quenching with water, the product was extracted with EtOAc and washed with 1 N HCl, concentrated *in vacuo*, triturated with MeOH, filtered and washed with MeOH. A mixture of the carboxylic acid and ester (469.0 mg) was obtained. The mixture of products were both committed to the hydrolysis conditions: 4 mL dioxane, 0.4 mL water, and 1 KOH pellet (~90 mg) were added to the mixture of acid and ester and shook at 50<sup>0</sup> C for 3 hrs. The hydrolysis was monitored by HPLC. The product was

dissolved in EtOAc and washed with 1 N HCl, concentrated *in vacuo*, triturated with MeOH, filtered, and washed with MeOH to obtain 246.8 mg (28%) of 5-bromo-2-({4-  
 5 [(5-fluoro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 11.95 (s, 1 H), 8.43 (d, *J* = 9 Hz, 1 H), 8.19 (d, *J* = 9 Hz, 2 H), 8.07 (d, *J* = 3 Hz, 1 H), 8.05 (d, *J* = 9 Hz, 2 H), 7.96 (dd, *J* = 9, 4 Hz, 1 H), 7.91 (d, *J* = 4 Hz, 1 H), 7.82 (dd, *J* = 9, 2 Hz, 1 H), 7.42 (dd, *J* = 9, 3 Hz, 1 H), 7.20 (td, *J* = 9, 3 Hz, 1 H), 6.86 (d, *J* = 4 Hz, 1 H); IR (drift) 1692, 1670, 1601, 1524 (s), 1462, 1388 (s), 1290, 1242, 1234, 1218 (s), 1181 (s), 1140 (s), 742, 649 (s), 607 (s), cm<sup>-1</sup>. MS (ESI-) for C<sub>22</sub>H<sub>14</sub>BrFN<sub>2</sub>O<sub>5</sub>S *m/z* 516.9 (M-H, Br isotope)<sup>-</sup>. HPLC [2] shows one major peak at  
 10 6.56 min (98%). Anal. Calcd for C<sub>22</sub>H<sub>14</sub>BrFN<sub>2</sub>O<sub>5</sub>S: C, 51.08; H, 2.73; N, 5.41; Br, 15.44; S, 6.20. Found: C, 51.05; H, 2.64; N, 5.39.

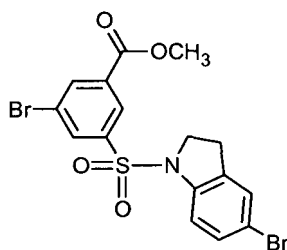
Other compounds were prepared by the above procedure making non-critical variations.

5-bromo-2-{{4-(1H-indol-1-ylsulfonyl)benzoyl}amino}benzoic acid, 5-bromo-2-({4-  
 15 [(6-fluoro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid, 5-bromo-2-({4-[(5-chloro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid, 5-bromo-2-({4-[(6-chloro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid, 5-bromo-2-({4-[(6-chloro-5-fluoro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid, 5-bromo-2-{{4-(1H-pyrrol-1-ylsulfonyl)benzoyl}amino}benzoic acid, 5-bromo-2-({4-[(5-methoxy-  
 20 1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid, 5-bromo-2-{{4-(1H-pyrrolo[2,3-b]pyridin-1-ylsulfonyl)benzoyl}amino}benzoic acid

### Scheme 1.3

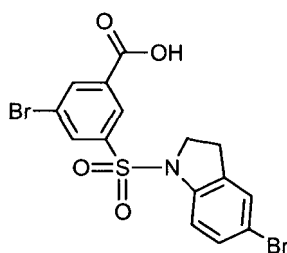


**Preparation of Methyl 3-bromo-5-[(5-bromo-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoate**



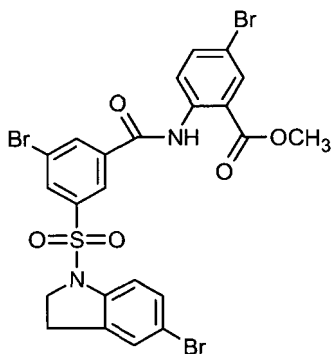
- 5 A solution of 5-bromoindoline (528 mg, 2.67 mmol, Lancaster) and triethylamine (650  $\mu$ L, 4.67 mmol) in  $\text{CH}_2\text{Cl}_2$  (8 mL) was added to a solution of methyl 3-bromo-5-(chlorosulfonyl)benzoate (737 mg, 2.35 mmol) in  $\text{CH}_2\text{Cl}_2$  (10 mL). The mixture was stirred overnight and then diluted to 100 mL with  $\text{CH}_2\text{Cl}_2$ . This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was
- 10 evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 50%  $\text{CH}_2\text{Cl}_2$ /heptane to 75%  $\text{CH}_2\text{Cl}_2$ /heptane as eluent. Yield was 945 mg of pale yellow solid.

- 15 **Preparation of 3-Bromo-5-[(5-bromo-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoic acid**



To a mixture of the corresponding methyl ester (841 mg, 1.77 mmol) in methanol (20 mL) was added 1 M aqueous sodium hydroxide (3.0 mL). The mixture was stirred in a 50 °C oil bath for 10 minutes and then at 60 °C for 15 minutes. The mixture was still a slurry, so 10 mL of dioxane was added. Heat was removed after an additional 40 minutes. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The organics were washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. They were then dried over MgSO<sub>4</sub> and evaporated yielding 807 mg of white solid.

**Methyl 5-bromo-2-({3-bromo-5-[(5-bromo-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoate**

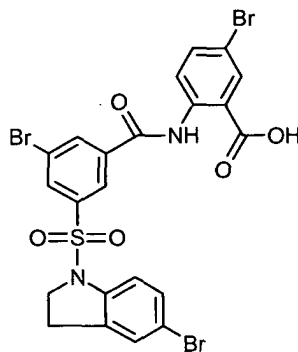


To 3-bromo-5-[(5-bromo-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoic acid (583 mg, 1.26 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (25 mL) was added DMF (20 µL) and oxalyl chloride (220 µL, 2.52 mmol). The mixture was stirred for 1 hour, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (10 mL), and methyl 2-amino-5-bromobenzoate (267 mg, 1.16 mmol, Avocado) in pyridine (4 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. Some THF was added to help solubility. This mixture was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The organics were evaporated, and the residue was dissolved in hot THF. This solution was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 50% CH<sub>2</sub>Cl<sub>2</sub>/heptane to 100% CH<sub>2</sub>Cl<sub>2</sub> as eluent. Yield was 603 mg of white solid.

**General Method D: (hydrolysis of the methyl ester)**

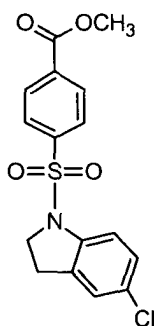


**5-Bromo-2-({3-bromo-5-[(5-bromo-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid**



To a mixture of the corresponding methyl ester (374 mg, 0.556 mmol) in dioxane (30 mL) was added 1 M aqueous sodium hydroxide (1.1 mL). The mixture was stirred in a 60 °C oil bath for 90 minutes. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The organics were washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. They were then dried over MgSO<sub>4</sub> and evaporated. The residue was recrystallized from hot ethanol/THF. The solids were washed with ethanol and then dried at 100 °C under vacuum yielding 266 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.14 (s, 1 H), 8.48 (d, *J* = 8.7 Hz, 1 H), 8.36 (s, 1 H), 8.31 (s, 1 H), 8.19 (s, 1 H), 8.12 (d, *J* = 2.0 Hz, 1 H), 7.86 (dd, *J* = 8.7, 2.5 Hz, 1 H), 7.39-7.49 (m, 3 H), 4.04 (t, *J* = 8.4 Hz, 2 H), 2.99 (t, *J* = 8.4 Hz, 2 H).

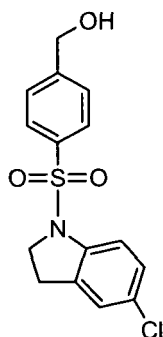
**Preparation of Methyl 4-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoate**



To 4-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoic acid (456 mg, 1.35 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (30 mL) was added DMF (15 μL) and oxalyl chloride (150 μL, 1.72 mmol). The mixture was stirred for 5 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub>

(10 mL). Methanol (2 mL) and pyridine (2 mL) in CH<sub>2</sub>Cl<sub>2</sub> (6 mL) were added. The mixture was stirred for 30 minutes and then added to a separatory funnel with 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. This solution was washed with 100 mL of 1 M aqueous HCl, 100 mL of saturated aqueous NaHCO<sub>3</sub>, another 100 mL of HCl, and 100 mL of brine. The  
5 CH<sub>2</sub>Cl<sub>2</sub> was dried over MgSO<sub>4</sub> and evaporated yielding 464 mg of white solid.

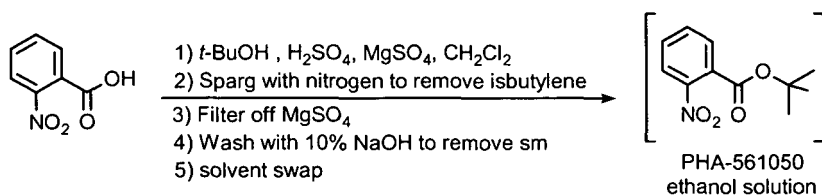
**Preparation of {4-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}methanol**



10

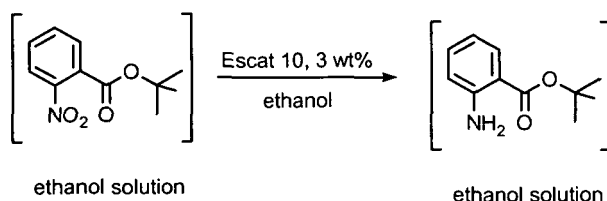
To a solution of methyl 4-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoate (396 mg, 1.13 mmol) in THF (20 mL) was added lithium borohydride (0.40 mL of 2.0 M solution in THF, 0.80 mmol, Aldrich). HPLC analysis after 1.5 hours indicated <10% reaction, so lithium aluminum hydride (0.60 mL of 1.0 M solution in THF,  
15 Aldrich) was added at -78 °C. The mixture was stirred at -78 °C for 15 minutes and then warmed to room temperature. The reaction was quenched by the addition of water (25 µL) followed by 6 M aqueous NaOH (25 µL) followed by another portion of water (75 µL). The mixture was filtered, and the filtrate was evaporated in the presence of silica gel. The product was purified by chromatography using a Biotage  
20 Flash 40 M silica cartridge with a gradient from CH<sub>2</sub>Cl<sub>2</sub> to 10% EtOAc in CH<sub>2</sub>Cl<sub>2</sub> as eluent. Yield was 290 mg of white solid.

**Preparation of *t*-butyl 2-nitrobenzoate**



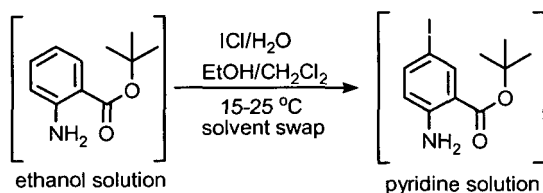
A 22 L round bottom flask, equipped with an mechanical stirrer, thermocouple, and a 1 L addition funnel, was charged with 500 g (2.99 moles, 1.0 equiv) of 2-nitrobenzoic acid (Avocado Research Chemicals Ltd, 98%) and 1.44 kg (11.97 moles, 4 equiv) of anhydrous magnesium sulfate (EM Science, 98%). To the solids were charged 12.5 L (25 mL/g) of CH<sub>2</sub>Cl<sub>2</sub> (EM Science, 99.96%) and 1.43 L (2.99 moles, 1.0 equiv) of *t*-butyl alcohol (Aldrich, 99 + % A.C.S. Reagent). The addition funnel was charged with 1.59 mL (2.99 moles, 1.0 equiv) of concentrated sulfuric acid (Mallinckrodt, 95.7%) and the entire system was sealed via use of a Teflon cap (loose fit; internal pressure does not exceed 11 psi; theory = 10.5 psi). The resulting suspension was cooled to 16 °C using a water bath and 159 mL (2.99 moles, 1.0 equiv) of concentrated sulfuric acid was added at a rate of 2.8 mL/min, maintaining an internal temperature less than 25 °C. The resulting off-white suspension was stirred at room temperature for 14 hours at which time the HPLC assay indicated the reaction was at 92% conversion. The suspension was sparged with nitrogen for 15 min using ½ inch ID Teflon tubing and filtered through a sintered glass funnel (course) with the aid of house vacuum (ca. 16 torr; filtration time of 1.0 h). The cake was rinsed with CH<sub>2</sub>Cl<sub>2</sub> (500 mL, 1 mL/g). The combined filtrates were charged to a 30 L wash tank and diluted with 2 L of water (pH = 1.0). To the resulting biphasic mixture was added 2.5 L of 10% NaOH over a 15 min period (8 °C exotherm; pH = 12.0). The resulting yellow-colored aqueous layers were separated from the clear, colorless organic layer. The organic layer was concentrated *in vacuo* at 16 torr using a 37 °C water bath to provide a 93% yield (621g, 2.78 moles) as a light yellow oil. To ensure removal of residual CH<sub>2</sub>Cl<sub>2</sub>, the oil was dissolved in 2 L of absolute ethanol (AAPER, 200 proof) and concentrated *in vacuo* at 16 torr using a 57 °C water bath. The potency of the material was determined to be 99.2% (GC) and 99.0% (HPLC) and was taken on directly to the next step without further purification.

### Preparation of *t*-butyl 2-aminobenzoate



Escat 10 catalyst (18.63 g, 3 wt%) was charged to the 10L autoclave followed by *t*-butyl nitrobenzoate (621g, 2.78 moles) in ethanol (7L). The vessel was sealed and purged three times with nitrogen (60 psig) and three times with hydrogen (60 psig). The vessel was then pressurized to 50 psig with hydrogen and allowed to run holding the exotherm at 40 °C through external cooling. The reaction was run until the hydrogen uptake stopped (45 minutes). The reaction was determined to be complete by both TLC and HPLC after 1 h and 10 min. The reaction was filtered through a 0.4  $\mu$  filter to remove the catalyst, and the catalyst cake was rinsed with ethanol (1.5 L). The product solution was then concentrated *in vacuo* at 16 torr using a 45 °C water bath to a volume of 1620 mL (3 mL/g) and taken on directly into the next step. An aliquot of the solution was concentrated and analyzed by both NMR and GC. The GC potency of the final product was 100%, and the NMR spectra were consistent with the structure of the title compound.

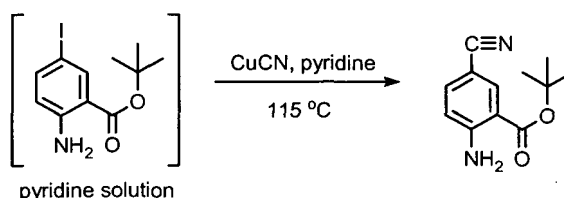
#### Preparation of *t*-butyl 2-amino-5-iodobenzoate



A 12 L round bottom flask, equipped with a thermocouple, nitrogen adapter and a 1 L addition funnel, was charged with a solution of *t*-butyl 2-aminobenzoate (537g, 2.78 moles; lot 36648-tjb-40) in ethanol (1620 ml, 3 ml/g). To this golden solution was added water (615.6 ml) resulting in a biphasic mixture. This mixture was cooled to between 15 and 20 °C with a cold-water bath. A 1.0 M solution of ICl in CH<sub>2</sub>Cl<sub>2</sub> (Aldrich lot #14127JO, 3.11 L, 3.11 moles, 1.12 equiv.) was charged in portions to the addition funnel and was added to the rapidly stirred mixture maintaining the temperature between 15 and 25 °C. The addition time was 2.25 hours and the temperature range observed was 16.5 to 20.4 °C. The resulting red brown mixture was stirred at room temperature for 1 hour at which time the GC assay showed the reaction was complete. The reaction was diluted with 920 mL of water and quenched with 456 mL of 38% aq. sodium bisulfite (Webb Chem lot #10464519) resulting in a slight exotherm to 24.0 °C. This mixture was stirred for 15 minutes before separating the

phases. The methylene chloride layer was combined with water (3.7L) and stirred for 15 minutes before separating the phases. A NaOH solution was prepared by diluting 10% NaOH (460ml) in water (2.3L). To the methylene chloride layer was added this dilute NaOH solution (2.1L). The pH of the basic phase was 6.56. The phases were separated and the methylene chloride layer was concentrated to a low volume *in vacuo* at 16 torr using a bath temp of 45 °C. Pyridine (4L) was added, and the resulting solution was concentrated to ca. 1.0 mL/g *in vacuo* at 16 torr using a 62 °C water bath. The low volume pyridine/product mixture was diluted with pyridine to the target volume of 3.1L (3.5 mL/g). A sample (10mL) was concentrated removing the pyridine on the rotovap and high vacuum to yield 3.12 g of an orange brown solid of 96% potency by GC. GC assay of pyridine solution indicated that neither EtOH nor methylene chloride were present, so the solution was taken on directly into the next step.

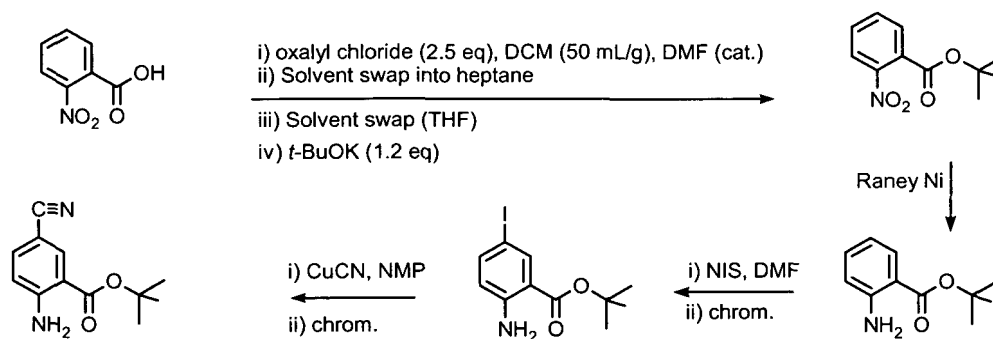
#### Preparation of *t*-butyl 2-amino-5-cyanobenzoate



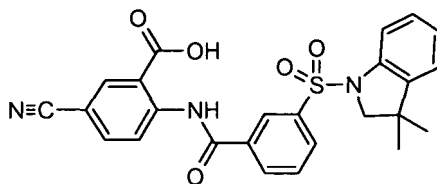
A 5 L Morton flask equipped with a mechanical stirrer (sturdy blade), thermocouple, and a reflux condenser was charged with 299g (3.34 moles, 1.2 equiv) of CuCN (Aldrich, 99%). To the slowly stirred CuCN was added a cool (10 °C) solution of *t*-butyl 2-amino-5-iodobenzoate (887g, 2.78 moles, 1.0 equiv) in pyridine (3.5 mL/g including the volume occupied by *t*-butyl 2-amino-5-iodobenzoate). The resulting orange suspension was heated to 115 °C over 45 min to produce a black solution. The solution was maintained at 115 °C for 14 h at which point GC indicated the reaction was complete. The solution was cooled to 90 °C and transferred by ½ inch ID Teflon cannula to a stirred suspension of solka floc (powdered cellulose, 460 g) in 14 L of methyl-*tert*-butyl ether (EM Science, 99.95%) at 2 °C, maintaining an internal temperature less than 13 °C. The resulting yellow-green suspension was filtered through a sintered glass frit (course frit, 16 torr vacuum) and the cake was rinsed with 4 L of MTBE (EM Science, 99.95%). The filtrate was washed (1 x 8 L H<sub>2</sub>O, 3 x 2 L

of 10% NH<sub>4</sub>OH in 23% NH<sub>4</sub>Cl), and the organics were concentrated *in vacuo* at 16 torr using a 50 °C water bath to a volume of 3 L (3.4 mL/g). The solution was split in half and crystallized in two portions. One half of the solution was charged to a 22 L flask containing heptanes (8L). The flask was set up for atmospheric distillation and heptanes (4L) was added to bring total volume of heptanes to 12 L. The mixture was distilled atmospherically to remove 4 L of distillate (pot temp of 98 °C; head temp of 96 °C). The pot was charged with 4 L of heptanes, and another 4 L of distillate was removed. A second 4 L charge of heptanes was made and 2.4 L of distillate was removed via atmospheric distillation; thus reducing the pot volume to 8.9 L (20mL/g). GC assay of the final distillate indicated the following volume percent ratios of pyridine and MTBE, respectively: 2.08% and 1.51 %. The heating mantle was removed, and the solution was cooled to induce crystallization (crystal formation was first noted at about 56 °C). The slurry was stirred at room temperature for 4 h, and the solids were isolated by vacuum filtration on a 3L frit. The cake was slurry washed with room temperature heptanes (2 x 500 ml) and dried on a nitrogen press to produce 224.2 g of an off-white solid (GC potency of 100%). Crystallization of the second half of the material produced another 241 g; thus a 70% yield from 2-nitrobenzoic acid was achieved.

An alternative methodology for producing t-butyl 2-amino-5-cyanobenzoate is shown below.

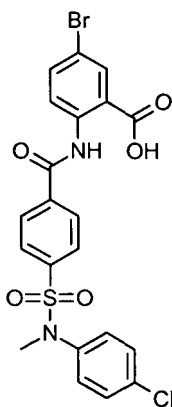


**5-Cyano-2-({3-[(3,3-dimethyl-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid**



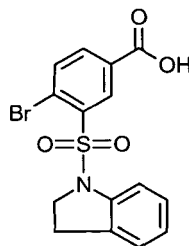
To a solution of 3-(chlorosulfonyl)benzoic acid (456 mg, 2.07 mmol, Aldrich) in  $\text{CH}_2\text{Cl}_2$  (15 mL) was added DMF (15  $\mu\text{L}$ ) followed by oxalyl chloride (270  $\mu\text{L}$ , 3.10 mmol). After stirring for 1.5 hours, the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in toluene (15 mL), and methyl 2-amino-5-cyanobenzoate (370 mg, 2.10 mmol) was added. The mixture was heated in a 105 °C oil bath for 2 hours, and the toluene was then removed by rotary evaporation. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (6 mL), and a mixture of 3,3-dimethylindoline, described by Kucerovy et al. in *Synth. Commun.* **1992**, 22(5), 729-733, (342 mg, 2.32 mmol) and triethylamine (600  $\mu\text{L}$ , 4.31 mmol) in  $\text{CH}_2\text{Cl}_2$  (6 mL) was added. This mixture was stirred overnight and then added to a separatory funnel with 100 mL of  $\text{CH}_2\text{Cl}_2$ . This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from  $\text{CH}_2\text{Cl}_2$  to 1% EtOAc in  $\text{CH}_2\text{Cl}_2$  as eluent. Yield was 728 mg of white solid as the methyl ester. The methyl ester was hydrolyzed according to method D yielding 292 mg of white solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.57 (s, 1 H), 8.80 (d,  $J$  = 8.7 Hz, 1 H), 8.41-8.44 (m, 2 H), 8.24 (d,  $J$  = 7.9 Hz, 1 H), 8.09-8.14 (m, 2 H), 7.83 (t,  $J$  = 7.9 Hz, 1 H), 7.55 (d,  $J$  = 8.1 Hz, 1 H), 7.24 (t,  $J$  = 7.7 Hz, 1 H), 7.18 (d,  $J$  = 7.7 Hz, 1 H), 7.02 (t,  $J$  = 7.5 Hz, 1 H), 3.73 (s, 2 H), 1.08 (s, 6 H).

**5-Bromo-2-[(4-[(4-chlorophenyl)(methyl)amino]sulfonyl}benzoyl)amino]benzoic acid**



Dimethyl formamide (15  $\mu$ L) and oxalyl chloride (1.5 mL, 17 mmol) were added sequentially to a mixture of 4-[[[(4-chlorophenyl)(methyl)amino]sulfonyl}benzoic acid (2.82 g, 8.66 mmol) in  $\text{CH}_2\text{Cl}_2$  (60 mL). The resulting solution was stirred for 3 hours after which the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (50 mL), and methyl 2-amino-5-bromobenzoate (1.83 g, 7.95 mmol, Avocado) in pyridine (15 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 150 mL of  $\text{CH}_2\text{Cl}_2$ . The resulting solution was washed with 2 X 100 mL of 1M aqueous HCl and 100 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 s silica cartridge with  $\text{CH}_2\text{Cl}_2$  as the eluent. Product was isolated as 3.73 g (87%) of a white solid as the methyl ester. The methyl ester was hydrolyzed according to method B.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.12 (s, 1 H), 8.56 (d,  $J$  = 8.7 Hz, 1 H), 8.10-8.14 (m, 3 H), 7.88 (dd,  $J$  = 8.7, 2.5 Hz, 1 H), 7.74 (d,  $J$  = 8.1 Hz, 2 H), 7.43 (d,  $J$  = 8.7 Hz, 2 H), 7.18 (d,  $J$  = 8.7 Hz, 2 H), 3.18 (s, 3 H).

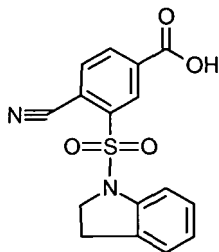
#### Preparation of 4-Bromo-3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoic acid



A solution of indoline (4.1 g, 34 mmol, Aldrich) and triethylamine (7.0 mL, 50 mmol) in methanol (20 mL) was added by cannula to solid 4-bromo-3-(chlorosulfonyl)benzoic acid (7.30 g, 24.4 mmol) with stirring in an ice bath. The mixture was allowed to warm slowly to room temperature and stirred overnight. It was added to a separatory funnel with 80 mL of aqueous 1 M NaOH, and this solution was washed with 2 X 100 mL of  $\text{CH}_2\text{Cl}_2$ . The aqueous layer was then acidified with concentrated HCl. The precipitate was washed with water followed by heptane and then recrystallized from toluene/ethanol. The crystals were washed with toluene followed by heptane and then dried at 100  $^{\circ}\text{C}$  under vacuum yielding 2.75 g of white solid. A second crop of 1.39 g of tan solid was also collected.



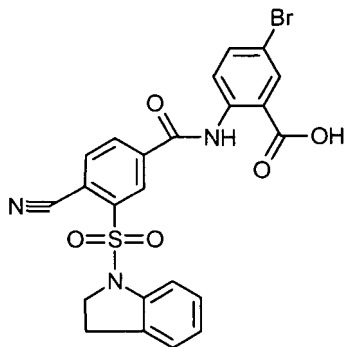
### Preparation of 4-Cyano-3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoic acid



5 A mixture of copper (I) cyanide (755 mg, 8.43 mmol) and 4-bromo-3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoic acid (2.05 g, 5.36 mmol) in NMP (15 mL) was heated to 160 °C under nitrogen for 1 hour. The mixture was added to a flask with 150 mL of EtOAc and 100 mL of water and stirred for 30 minutes. It was then filtered through a plug of celite. The phases were separated, and the water was extracted with an  
10 additional 2 X 100 mL of EtOAc. The combined EtOAc was washed with 3 X 100 mL of water and dried over MgSO<sub>4</sub>. The solvent was removed, and the brown residue was recrystallized from hot ethanol. The crystals were washed with methanol followed by heptane and then dried at 100 °C under vacuum. Yield was 1.25 g of tan solid.

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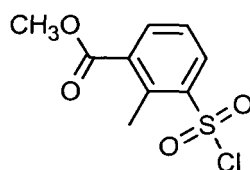
### 5-Bromo-2-{{4-cyano-3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl}amino}benzoic acid



To 4-cyano-3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoic acid (1.22 g, 3.72 mmol) in  
20 CH<sub>2</sub>Cl<sub>2</sub> (30 mL) was added DMF (20 µL) and oxalyl chloride (650 µL, 7.45 mmol). The mixture was stirred for 2.3 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved as best as possible in CH<sub>2</sub>Cl<sub>2</sub> (30 mL), and methyl 2-amino-5-bromobenzoate ( 762 mg, 3.31 mmol, Avocado) in pyridine (15 mL) was added. The mixture was stirred overnight and then

added to a separatory funnel with 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The CH<sub>2</sub>Cl<sub>2</sub> was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with CH<sub>2</sub>Cl<sub>2</sub> as eluent. Yield was 1.31 g of yellow solid. The methyl ester was hydrolyzed according to Method D to yield 615 mg of yellow solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.24 (s, 1 H), 8.57 (s, 1 H), 8.51 (d, *J* = 8.7 Hz, 1 H), 8.37 (d, *J* = 7.6 Hz, 1 H), 8.32 (d, *J* = 7.6 Hz, 1 H), 8.14 (d, *J* = 2.5 Hz, 1 H), 7.88 (dd, *J* = 8.9, 2.3 Hz, 1 H), 7.43 (d, *J* = 8.1 Hz, 1 H), 7.16-7.24 (m, 2 H), 7.01 (t, *J* = 7.6 Hz, 1 H), 4.20 (t, *J* = 8.4 Hz, 2 H), 3.05 (t, *J* = 8.4 Hz, 2 H).

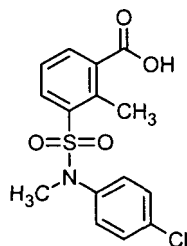
### Preparation of Methyl 3-(chlorosulfonyl)-2-methylbenzoate



A flask was charged with methyl 2-methyl-3-nitrobenzoate (Aldrich, 5.0 g, 25.6 mmol) and tin (II) chloride dihydrate (28.9 g, 128 mmol, 5.0 eq). The solids were suspended in EtOAc (80 mL), and upon heating to reflux under N<sub>2</sub> the solids completely dissolved. After two hours the cooled reaction was poured into 350 mL EtOAc and then washed 4x with 1.0M NaOH, 1x with water and 1x with brine (350 mL each). The organic layer was dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and the solvent evaporated. The resultant crude oil (2.9 g) was suspended in 60 mL of a 2:1 solution of concentrated HCl and glacial acetic acid. The reaction was cooled to -10 °C and a solution of sodium nitrite (1.33g, 19.34 mmol) in 3.0 mL water was added drop wise over stirring at a rate that maintained the internal reaction temperature below -5 °C. The reaction became an orange solution as the SM slowly dissolved. In a separate flask, copper (I) chloride (435 mg, 25 mol%) was suspended in 30 mL of a saturated (30% w/w) solution of sulfur dioxide gas in glacial acetic acid. The mixture was cooled on an ice bath over stirring, and after 2.5 hours the diazonium solution was added portion wise to the copper mixture over 15 minutes. The addition evolved gas and produced a lime green solution, which came to RT and was stirred overnight. The reaction was poured into ice water (200 mL) to afford an oil at the bottom of a pale blue solution. The solution was extracted 2x with CH<sub>2</sub>Cl<sub>2</sub> (150 mL ea) and the organic phase was washed 2x with saturated NaHCO<sub>3</sub> and brine (250 mL ea). The

golden organic solution was dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and the solvent evaporated. The crude residue was purified on a Biotage Flash 40M+ (100g) silica cartridge using a gradient of 20% heptane in CH<sub>2</sub>Cl<sub>2</sub> to 100% CH<sub>2</sub>Cl<sub>2</sub>. The combined fractions were evaporated and the product was dried under high vacuum at RT to afford 2.2 g of pale pink solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 7.96 (dd, *J* = 7.7, 1.5 Hz, 1 H), 7.59 (dd, *J* = 7.7, 1.5 Hz, 1 H), 7.23 (t, *J* = 7.7 Hz, 1 H), 3.82 (s, 3 H), 2.56 (s, 3 H).

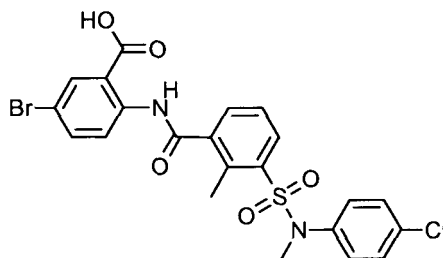
### Preparation of 3-[[4-(chlorophenyl)(methyl)amino]sulfonyl]-2-methylbenzoate



Methyl 3-(chlorosulfonyl)-2-methylbenzoate, (494 mg, 1.99 mmol) was taken up in dry CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and treated with 4-chloro-N-methylaniline (1.01 mL, 8.35 mmol, Aldrich) in dry pyridine (15 mL). The bright yellow solution was heated to 75 °C. After one hour HPLC indicated the reaction was complete and the mixture was poured into EtOAc (125 mL). The organic phase was washed 3x with 1.0M HCl, 1x with saturated NaHCO<sub>3</sub> and 1x with brine (100 mL each). After drying over Na<sub>2</sub>SO<sub>4</sub> the solution was filtered and the solvent was evaporated to afford an amber oil, which was purified on a Biotage Flash 40M+ (100g) silica cartridge using a linear gradient of 35% to 5% heptane in CH<sub>2</sub>Cl<sub>2</sub>. The solvent was evaporated from the product fractions and the product was dried under high vacuum at RT to afford 637 mg (90%) of a colorless oil. 508 mg, 1.44 mmol of the oil was dissolved in MeOH (15 mL) and treated with 1.0M LiOH (3.0 mL, 3.0 mmol). After stirring at 40 °C for 1 hour and then overnight at RT, the reaction was complete by HPLC and OAMS showed the correct *m/z* for product. The reaction was poured into 1.0M HCl (100 mL), and the white precipitate was extracted into EtOAc (150 mL). The organic layer was then 1x with 1.0M HCl and 1x with brine (125 mL each). The organic layer was dried over MgSO<sub>4</sub>, filtered and evaporated to dryness. The resultant product was dried under vacuum at 100 °C overnight to afford 461 mg (94%) of off-white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 13.41 (br s, 1 H), 7.94 (d, *J* = 3.3 Hz, 1 H), 7.92 (d, *J* = 3.1

Hz, 1H), 7.49 (t,  $J = 7.9$  Hz, 1 H), 7.39-7.47 (m, 2 H), 7.22-7.31 (m, 2 H), 3.21 (s, 3 H), 2.45 (s, 3 H).

**5-Bromo-2-[(3-[[[(4-chlorophenyl)(methyl)amino]sulfonyl]-2-methylbenzoyl)amino]benzoic acid**

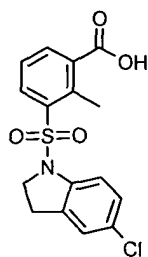


3-[[[(4-chlorophenyl)(methyl)amino]sulfonyl]-2-methylbenzoate (404 mg, 1.19 mmol) was suspended in dry  $\text{CH}_2\text{Cl}_2$  (10 mL) and DMF (10  $\mu\text{L}$ ) under  $\text{N}_2$ . The solution was treated with oxalyl chloride (Aldrich, 0.192 mL, 2.2 mmol) and stirred while gas evolved. After one hour the excess solvent and oxalyl chloride were evaporated and the resultant residue was taken up in dry  $\text{CH}_2\text{Cl}_2$  (10 mL). Methyl-2-amino-5-bromobenzoate (Aldrich, 230 mg, 1.0 mmol) was added as a solution in pyridine (3 mL) and the amber solution stirred at RT. After 2 hours HPLC indicated the reaction was complete. The mixture was diluted with  $\text{CH}_2\text{Cl}_2$  (100 mL) and washed 2x with 1.0M HCl followed by brine (100 mL each). The organic layer was evaporated and purified on a Biotage Flash 25M+ (40 g) silica cartridge using  $\text{CH}_2\text{Cl}_2$ . The combined fractions were evaporated and the product was dried under vacuum at 100  $^\circ\text{C}$  to afford 535mg (97%) of a glass-like solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  10.88 (s, 1 H), 8.05 (d,  $J = 8.9$  Hz, 1 H), 7.99 (d,  $J = 2.3$  Hz, 1 H), 7.93 (D,  $J = 7.5$  Hz, 1 H), 7.86 (dd,  $J = 8.8, 2.4$  Hz, 1 H), 7.80 (d,  $J = 7.3$  Hz, 1 H), 7.57 (t,  $J = 7.9$  Hz, 1 H), 7.45 (d,  $J = 8.7$  Hz, 2 H), 7.29 (d,  $J = 8.7$  Hz, 2 H), 3.83 (s, 3 H), 3.24 (s, 3 H), 2.39 (s, 3 H). 322 mg of the methyl ester solid was dissolved in hot dioxane (10 mL), and after cooling was treated with 1.0M LiOH (1.0 mL, 1.0 mmol). After stirring overnight at RT the reaction was complete by HPLC and OAMS showed correct  $m/z$  for the product. The solvent was evaporated and the residue was poured into 1.0M HCl (100 mL) to afford a white precipitate. The product was extracted into EtOAc (125 mL) and washed 3x with 1.0M HCl, and 1x with brine (100 mL each). The organic layer was dried over  $\text{Na}_2\text{SO}_4$ , filtered and evaporated to dryness. The crude product was recrystallized from hot MeOH/EtOH. The resultant product was dried at 100  $^\circ\text{C}$  under

vacuum to afford 213 mg (68%) of white crystals.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  11.35 (s, 1 H), 8.39 (d,  $J = 8.9$  Hz, 1 H), 8.07 (d,  $J = 2.5$  Hz, 1 H), 7.92 (dd,  $J = 8.1$ , 1.0 Hz, 1 H), 7.81-7.89 (m, 2 H), 7.56 (t,  $J = 7.8$  Hz, 1 H), 7.41-7.48 (m, 2 H), 7.24-7.34 (m, 2 H), 3.23 (s, 3 H), 2.39 (s, 3 H).

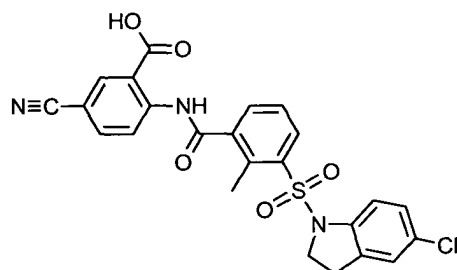
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### Preparation of 3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]-2-methylbenzoic acid



Methyl 3-(chlorosulfonyl)-2-methylbenzoate, (673 mg, 2.71 mmol) was taken up in  
10 dry  $\text{CH}_2\text{Cl}_2$  (5 mL) and dry pyridine (5 mL). The golden solution was cooled to  $-10$   
 $^\circ\text{C}$  and treated with 5-chloroindoline (1.01 mL, 8.35 mmol, Aldrich) in dry  $\text{CH}_2\text{Cl}_2$  (5  
mL) to afford an intensely red-orange solution. A precipitate formed as the reaction  
warmed to RT. After one hour HPLC indicated the reaction was complete and the  
mixture was diluted to 150 mL with  $\text{CH}_2\text{Cl}_2$ . The organic phase was washed 1x with  
15 1.0M HCl, 1x with 1.0M NaOH, 1x with 1.0M HCl and 1x with brine (125 mL each).  
After drying over  $\text{Na}_2\text{SO}_4$  the solution was filtered and the solvent was evaporated.  
The resultant product was dried under high vacuum at RT to afford 900 mg (90%) of a  
peach colored oil. 780mg (2.13 mmol) of the oil was dissolved in MeOH (15 mL) and  
treated with 1.0M LiOH (5.0 mL, 5.0 mmol). After stirring at  $40$   $^\circ\text{C}$  for 1 hour and  
20 then overnight at RT, the reaction was complete by HPLC and OAMS showed the  
correct  $m/z$  for product. The reaction was poured into 1.0M HCl (125 mL), and the  
yellowish precipitate was extracted into EtOAc (150 mL). The organic layer was then  
2x with 1.0M HCl, 1x with water and 1x with brine (125 mL each). The organic layer  
was dried over  $\text{MgSO}_4$ , filtered and evaporated to dryness. The resultant product was  
25 dried under vacuum at  $100$   $^\circ\text{C}$  overnight to afford 711 mg (95%) of pinkish-orange  
solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  13.46 (br s, 1 H), 7.98 (d,  $J = 8.1$  Hz, 1 H),  
7.93 (d,  $J = 7.7$  Hz, 1 H), 7.50 (t,  $J = 7.9$  Hz, 1 H), 7.34 (d,  $J = 1.7$  Hz, 1 H), 7.19 (dd,  
 $J = 8.5$ , 2.1 Hz, 1 H), 7.09 (d,  $J = 8.5$  Hz, 1 H), 4.05 (t,  $J = 8.5$  Hz, 2 H), 3.12 (t,  $J =$   
8.5 Hz, 2 H), 2.66 (s, 3 H).

**2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]-2-methylbenzoyl}amino)-5-cyanobenzoic acid**



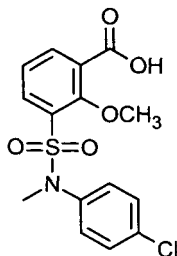
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3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]-2-methylbenzoic acid (553 mg, 01.57 mmol) was suspended in dry CH<sub>2</sub>Cl<sub>2</sub> (15 mL) and DMF (10 μL) under N<sub>2</sub>. The solution was treated with oxalyl chloride (0.274 mL, 3.14 mmol, Aldrich) and stirred while gas evolved. The reaction became homogenous and after one hour the excess solvent and oxalyl chloride was evaporated and the resultant residue was taken up in dry CH<sub>2</sub>Cl<sub>2</sub> (10 mL). Methyl-2-amino-5-cyanobenzoate (PHA-522499, 264 mg, 1.5 mmol) was added as a solution in pyridine (4 mL) and the amber solution stirred at RT. After 2.5 days HPLC indicated the reaction was nearly complete. After briefly boiling the reaction and cooling, the mixture was diluted to 150 mL with CH<sub>2</sub>Cl<sub>2</sub> and washed 2x with 1.0M HCl followed by brine (125 mL each). The organic layer was dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and evaporated. The resultant crude product was purified on a Biotage Flash 25M+ (40 g) silica cartridge using a linear gradient of 0-2% EtOAc in CH<sub>2</sub>Cl<sub>2</sub>. The resultant product still contained a small amount of residual cyanoanthranilate. The combined fractions were evaporated and the product was purified a second time on a Biotage Flash 40M+ (100 g) silica cartridge using 100% CH<sub>2</sub>Cl<sub>2</sub>. The combined fractions were evaporated and dried under high vacuum at RT to afford 594mg (77%) of an off-white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 11.21 (s, 1 H), 8.29 (d, *J* = 1.9 Hz, 1 H), 8.26 (d, *J* = 8.7 Hz, 1 H), 8.11 (dd, *J* = 8.6, 2.0 Hz, 1 H), 7.99 (dd, *J* = 8.1, 1.0 Hz, 1 H), 7.84 (dd, *J* = 7.7, 1.0 Hz, 1 H), 7.60 (t, *J* = 7.9 Hz, 1 H), 7.36 (d, *J* = 1.7 Hz, 1 H), 7.21 (dd, *J* = 8.7, 2.3 Hz, 1 H), 7.16 (d, *J* = 8.7 Hz, 1 H), 4.09 (t, *J* = 8.6 Hz, 2 H), 3.84 (s, 3 H), 3.15 (t, *J* = 8.4 Hz, 2 H), 2.61 (s, 3 H). The methyl ester was hydrolyzed as described above to afford 300 mg (77%) of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 11.73 (s, 1 H), 8.62 (d, *J* = 8.7 Hz, 1 H), 8.36 (d, *J* = 2.1 Hz, 1 H), 8.10 (dd, *J* = 8.7, 2.1 Hz, 1 H), 7.97 (d, *J* = 8.1 Hz, 1 H), 7.90 (d,

25

$J = 6.8$  Hz, 1 H), 7.57 (t,  $J = 7.9$  Hz, 1 H), 7.37 (s, 1 H), 7.20 (dd,  $J = 8.7, 2.1$  Hz, 1 H), 7.16 (d,  $J = 8.5$  Hz, 1 H), 4.07 (t,  $J = 8.6$  Hz, 2 H), 3.15 (t,  $J = 8.5$  Hz, 2 H), 2.62 (s, 3 H).

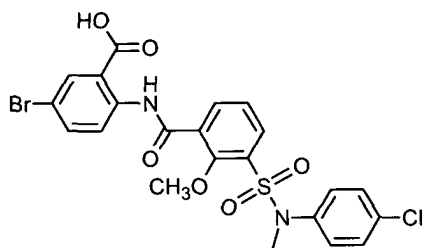
5 **Preparation of 3-[[[4-Chlorophenyl](methyl)amino]sulfonyl]-2-methoxybenzoic acid**



Methyl 3-amino-2-methoxybenzoate (1.27 g, 6.72 mmol) was dissolved in 30 mL of a 2:1 solution of concentrated HCl and glacial acetic acid. The reaction was cooled to –  
10 °C and a solution of sodium nitrite (696 mg, 10.1 mmol) in 3.0 mL water was added drop wise over stirring at a rate that maintained the internal reaction temperature below –5 °C. The reaction became a cloudy yellow-orange suspension. In a separate flask, copper (I) chloride (166 mg, 25 mol%) was suspended in 30 mL of a saturated (30% w/w) solution of sulfur dioxide gas in glacial acetic acid. The mixture was cooled on an ice bath over stirring, and after 30 minutes diazonium solution was added portion wise to the copper mixture over 15 minutes. The addition evolved gas and produced a dark green solution. The reaction was warmed to RT and was stirred for 3 hours with sulfur dioxide bubbling into the solution. The reaction was poured into ice water (200 mL) to afford a fine white precipitate in a pale blue solution. The solution was extracted 3x with EtOAc (150 mL ea) and the organic phase was neutralized by washing 3x with saturated NaHCO<sub>3</sub> (300 mL ea). The organic phase was then washed 2x with water and 1x with brine (250 mL ea). The golden organic solution was dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and the solvent evaporated. The crude residue was dried under high vacuum to afford a dark red oil. The oil was taken up in pyridine (15 mL) and treated with 4-chloro-N-methylaniline (0.280 mL, 2.3 mmol, Aldrich). The amber solution was heated stirred at RT, and after one hour HPLC indicated the reaction was complete. The mixture was diluted to 150 mL with DCM and then washed 2x with 1.0M HCl, 1x with 1.0M NaOH and 1x with brine (125 mL each). The solvent was evaporated to afford an amber oil, which was

purified on a Biotage Flash 40M (90g) silica cartridge using a linear gradient of 0 to 0.75% EtOAc in CH<sub>2</sub>Cl<sub>2</sub>. The solvent was evaporated from the product fractions and the product was dried under high vacuum at RT to afford 614 mg (72%) of a straw colored oil as the methyl ester. The methyl ester was hydrolyzed as described above to afford 544 mg (97%) of peach colored solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 13.50 (s, 1 H), 7.99 (dd, *J* = 7.7, 1.9 Hz, 1 H), 7.80 (dd, *J* = 7.9, 1.7 Hz, 1 H), 7.36-7.42 (m, 2 H), 7.30 (t, *J* = 7.9 Hz, 1 H), 7.19-7.26 (m, 2 H), 3.83 (s, 3 H), 3.32 (s, 3 H).

**5-Bromo-2-[(3-[(4-chlorophenyl)(methyl)amino]sulfonyl]-2-methoxybenzoyl)amino]benzoic acid**

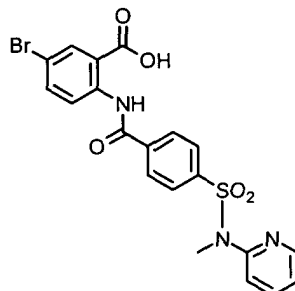


3-[(4-Chlorophenyl)(methyl)amino]sulfonyl}-2-methoxybenzoic acid (PHA-733277, 474 mg, 0.133 mmol) was dissolved in dry CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and DMF (25 μL) under N<sub>2</sub>. The solution was treated with oxalyl chloride (0.232 mL, 2.66 mmol, Aldrich) and stirred while gas evolved. The reaction was stirred at RT and after one hour the excess solvent and oxalyl chloride was evaporated and the resultant residue was taken up in dry CH<sub>2</sub>Cl<sub>2</sub> (10 mL). Methyl-2-amino-5-bromobenzoate (288 mg, 1.25 mmol, Avocado) was added as a solution in pyridine (3 mL) and the amber solution stirred at RT. After 90 minutes HPLC indicated the reaction was complete. The mixture was diluted to 150 mL with CH<sub>2</sub>Cl<sub>2</sub> and washed 2x with 1.0M HCl followed by brine (100 mL each). The organic layer was dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and evaporated. The resultant crude product was purified on a Biotage Flash 40M (90 g) silica cartridge using CH<sub>2</sub>Cl<sub>2</sub>. The combined fractions were evaporated and dried under vacuum at 100 °C to afford 530mg (72%) of an off-white solid as the methyl ester. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 11.52 (s, 1 H), 8.48 (d, *J* = 8.7 Hz, 1 H), 8.10 (d, *J* = 2.5 Hz, 1 H), 7.98 (dd, *J* = 7.8, 1.8 Hz, 1 H), 7.91 (dd, *J* = 8.9, 2.5 Hz, 1 H), 7.81 (dd, *J* = 7.9, 1.7 Hz, 1 H), 7.32-7.43 (m, 5 H), 3.89 (s, 3 H), 3.82 (s, 3 H), 3.40 (s, 3 H). The corresponding methyl ester was hydrolyzed as described above to afford a white solid.



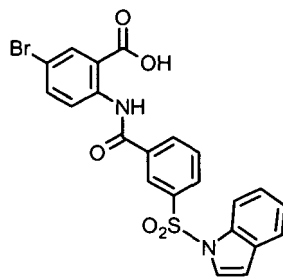
<sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.02 (s, 1 H), 8.70 (d, *J* = 9.1 Hz, 1 H), 8.14 (d, *J* = 2.5 Hz, 1 H), 8.01 (dd, *J* = 7.7, 1.2 Hz, 1 H), 7.90 (dd, *J* = 9.0, 2.4 Hz, 1 H), 7.76 (dd, *J* = 7.9, 1.5 Hz, 1 H), 7.11-7.44 (m, 5 H), 3.81 (s, 3 H), 3.39 (s, 3 H).

5 **5-bromo-2-[(4-{[methyl(pyridin-2-yl)amino]sulfonyl}benzoyl)amino]benzoic acid**



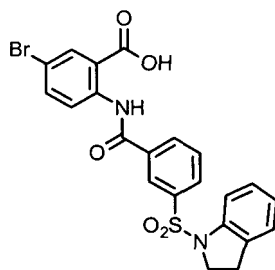
4-{[methyl(pyridin-2-yl)amino]sulfonyl}benzoic acid (292 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 0.5 mL, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (20% EtOAc in hexane) to provide 317 mg of the desired methyl ester (63%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (281 mg, 91%, 57% overall) was obtained as a tan solid after recrystallization from MeOH. H NMR (400 MHz, DMSO) 3.72 (s, 3H), 7.28 (dd, 1H), 7.56 (d, 1H), 7.81-7.91 (m, 4H), 8.07 (d, 2H), 8.12 (d, 1H), 8.32 (dd, 1H), 8.54 (d, 1H), 12.10 (s, 1H). C NMR (100 MHz, DMSO) 36.10, 101.83, 115.38, 120.21, 120.30, 122.15, 122.90, 128.33, 128.42, 133.62, 136.95, 138.77, 139.91, 140.30, 148.52, 153.13, 163.81, 168.81. MS (FAB) *m/z* (rel. intensity) 490 (MH<sup>+</sup>, 30), 492 (32), 490 (30), 414 (28), 413 (83), 109 (31), 107 (36), 95 (25), 91 (99), 57 (73), 55 (28). HRMS (FAB) calcd for C<sub>20</sub>H<sub>16</sub>BrN<sub>3</sub>O<sub>5</sub>S +H<sub>1</sub>, 490.0073, found 490.0067.

**5-bromo-2-{[3-(1H-indol-1-ylsulfonyl)benzoyl]amino}benzoic acid**



3-(1H-indol-1-ylsulfonyl)benzoic acid (301 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 0.5 mL, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide 287 mg of the desired methyl ester (56%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (53 mg, 11%, 6% overall) was obtained as a white solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 6.90 (d, 1H), 7.27 (t, 1H), 7.37 (t, 1H), 7.62 (d, 1H), 7.82 (t, 1H), 7.87-7.89 (m, 2H), 8.00 (d, 1H), 8.05 (d, 1H), 8.19-8.25 (m, 3H), 8.47 (s, 1H), 11.35 (s, 1H).

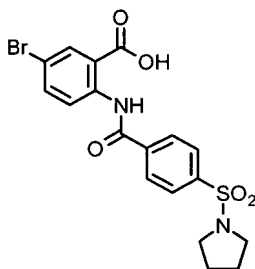
#### 5-bromo-2-[[3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl]amino]benzoic acid



3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoic acid (305 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 0.5 mL, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol)

was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide 381 mg of the desired methyl ester (74%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (344 mg, 93%, 68% overall) was obtained as a white solid after recrystallization from MeOH. H NMR (400 MHz, DMSO) 2.94 (t, 2H), 4.00 (t, 2H), 6.99 (t, 1H), 7.15-7.23 (m, 2H), 7.52 (d, 1H), 7.80 (t, 1H), 7.89 (dd, 1H), 8.05-8.07 (m, 2H), 8.20 (d, 1H), 8.28 (d, 1H), 8.35 (s, 1H), 11.40 (s, 1H).

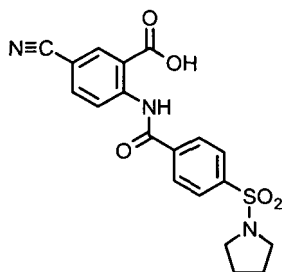
**5-bromo-2-[[4-(pyrrolidin-1-ylsulfonyl)benzoyl]amino]benzoic acid**



4-(pyrrolidin-1-ylsulfonyl)benzoic acid (255 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 0.5 mL, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide 331 mg of the desired methyl ester (71%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (308 mg, 96%, 68% overall) was obtained as a pale yellow solid

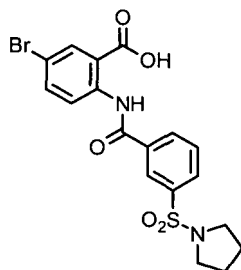
after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 1.67 (m, 4H), 3.19 (m, 4H), 7.88 (dd, 1H), 8.02 (d, 2H), 8.12-8.16 (m, 3H), 8.58 (d, 1H), 12.10 (s, 1H).

**5-cyano-2-{{4-(pyrrolidin-1-ylsulfonyl)benzoyl}amino}benzoic acid**



5 4-(pyrrolidin-1-ylsulfonyl)benzoic acid (255 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 0.5 mL, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL).  
10 Methyl 2-amino-5-cyanobenzoate (176 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to  
15 provide 293 mg of the desired methyl ester (71%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (262 mg, 92%, 65% overall) was obtained as a pale yellow solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 1.67 (m, 4H), 3.20  
20 (m, 4H), 8.04 (d, 2H), 8.11-8.18 (m, 3H), 8.42 (d, 1H), 8.80 (d, 1H), 12.25 (s, 1H).

**5-bromo-2-{{3-(pyrrolidin-1-ylsulfonyl)benzoyl}amino}benzoic acid**

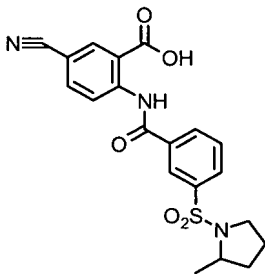


3-(pyrrolidin-1-ylsulfonyl)benzoic acid (255 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub>  
25 (10 mL) and (COCl)<sub>2</sub> added (725 mg, 0.5 mL, 5.7 mmol). A catalytic amount of

DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide 333 mg of the desired methyl ester (71%). The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over  $\text{Na}_2\text{SO}_4$  and then concentrated *in vacuo*. The title compound (309 mg, 96%, 68% overall) was obtained as a pale yellow solid after recrystallization from MeOH.  $^1\text{H}$  NMR (400 MHz, DMSO) 1.67 (m, 4H), 3.20 (m, 4H), 7.85-7.89 (m, 2H), 8.08 (d, 1H), 8.13 (d, 1H), 8.25 (d, 1H), 8.32 (s, 1H), 8.60 (d, 1H), 12.20 (s, 1H).

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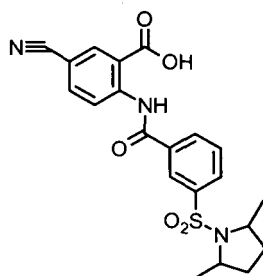
**5-cyano-2-({3-[(2-methylpyrrolidin-1-yl)sulfonyl]benzoyl}amino)benzoic acid**



3-[(2-methylpyrrolidin-1-yl)sulfonyl]benzoic acid (269 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-cyanobenzoate (176 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide 350 mg of the desired methyl ester (82%). The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over  $\text{Na}_2\text{SO}_4$  and then concentrated *in vacuo*.

The title compound (308 mg, 91%, 75% overall) was obtained as a white solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 1.25 (d, 3H), 1.41-1.47 (m, 2H), 1.59-1.67 (m, 1H), 1.77-1.83 (m, 1H), 3.12-3.18 (m, 1H), 3.36-3.42 (m, 1H), 3.69 (m, 1H), 7.88 (t, 1H), 8.12 (d, 1H), 8.13 (d, 1H), 8.25 (d, 1H), 8.34 (s, 1H), 8.42 (d, 1H), 8.83 (d, 1H), 12.55 (s, 1H).

**5-cyano-2-({3-[(2,5-dimethylpyrrolidin-1-yl)sulfonyl]benzoyl}amino)benzoic acid**



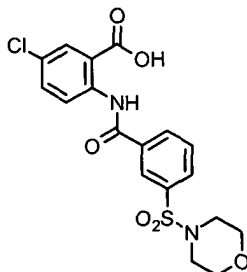
3-[(2,5-dimethylpyrrolidin-1-yl)sulfonyl]benzoic acid (283 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL). Methyl 2-amino-5-cyanobenzoate (176 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide 293 mg of the desired methyl ester (66%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (273 mg, 97%, 64% overall) was obtained as a tan solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 1.29 (d, 6H), 1.49 (m, 4H), 3.67 (m, 2H), 7.88 (t, 1H), 8.12 (d, 1H), 8.13 (d, 1H), 8.25 (d, 1H), 8.34 (s, 1H), 8.42 (d, 1H), 8.83 (d, 1H), 12.55 (s, 1H).

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**5-cyano-2-{{3-(pyrrolidin-1-ylsulfonyl)benzoyl}amino}benzoic acid** was produced from methyl 2-{{3-(chlorosulfonyl)benzoyl}amino}-5-cyanobenzoate. <sup>1</sup>H NMR (300 MHz, DMSO) 1.67 (m, 4H), 3.20 (m, 4H), 7.88 (t, 1H), 8.09-8.14 (m, 2H), 8.26 (d, 1H), 8.33 (s, 1H), 8.42 (d, 1H), 8.83 (d, 1H), 12.56 (s, 1H)

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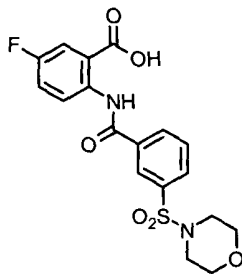
**5-chloro-2-[[3-(morpholin-4-ylsulfonyl)benzoyl]amino]benzoic acid**



3-(morpholin-4-ylsulfonyl)benzoic acid (271 mg, 1.0 mmol) was suspended in CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and (COCl)<sub>2</sub> added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in CHCl<sub>3</sub> (10 mL). Methyl 2-amino-5-chlorobenzoate (185 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (20% EtOAc in hexane) to provide 382 mg of the desired methyl ester (87%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (351 mg, 95%, 83% overall) was obtained as a white solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 2.93 (m, 4H), 3.65 (m, 4H), 7.77 (dd, 1H), 7.91 (t, 1H), 7.99-8.02 (m, 2H), 8.25-8.29 (m, 2H), 8.65 (d, 1H), 12.17 (s, 1H).

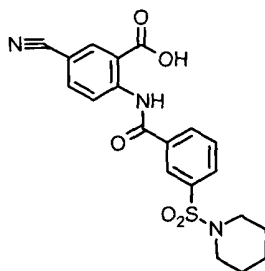
5-bromo-2-[[3-(morpholin-4-ylsulfonyl)benzoyl]amino]benzoic acid and 2-[[3-(morpholin-4-ylsulfonyl)benzoyl]amino]-5-nitrobenzoic acid were produced in a similar fashion utilizing appropriate starting materials.

**5-fluoro-2-[[3-(morpholin-4-ylsulfonyl)benzoyl]amino]benzoic acid**



3-(morpholin-4-ylsulfonyl)benzoic acid (271 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-fluorobenzoate (170 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (20% EtOAc in hexane) to provide 367 mg of the desired methyl ester (87%). The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over  $\text{Na}_2\text{SO}_4$  and then concentrated *in vacuo*. The title compound (328 mg, 92%, 80% overall) was obtained as a white solid after recrystallization from MeOH.  $^1\text{H}$  NMR (400 MHz, DMSO) 2.93 (m, 4H), 3.65 (m, 4H), 7.58 (m, 1H), 7.77 (dd, 1H), 7.90 (t, 1H), 8.00 (d, 1H), 8.26-8.29 (m, 2H), 8.60 (dd, 1H), 12.02 (s, 1H).

#### 5-cyano-2-([3-(piperidin-1-ylsulfonyl)benzoyl]amino)benzoic acid

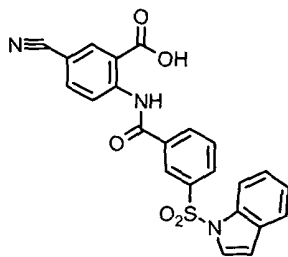


3-(piperidin-1-ylsulfonyl)benzoic acid (269 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-cyanobenzoate (176 mg, 1.0 mmol) was added followed by pyridine (1



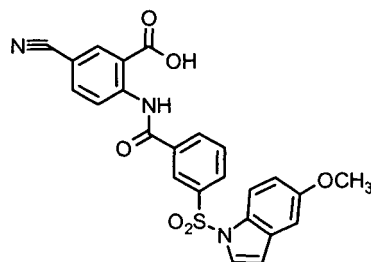
mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (10% EtOAc in hexane) to provide  
 5 307 mg of the desired methyl ester (72%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (279 mg, 94%) was obtained as a white solid after recrystallization from MeOH. H NMR (400 MHz, DMSO) 1.37 (m, 2H), 1.56 (m, 4H), 2.95 (m, 4H), 7.90  
 10 (t, 1H), 8.02 (d, 1H), 8.13 (dd, 1H), 8.27 (m, 2H), 8.42 (d, 1H), 8.83 (1H), 12.55 (s, 1H).

**5-cyano-2-([3-(1H-indol-1-ylsulfonyl)benzoyl]amino)benzoic acid**



15 Indole (150 mg, 1.25 mmol) was dissolved in 15 ml of THF. NaH (100 mg, 60% disp. in oil, 2.5 mmol) was added and resulting suspension stirred for 1 h. Methyl 2-([3-(chlorosulfonyl)benzoyl]amino)-5-cyanobenzoate (378 mg, 1.0 mmol) was then added and the reaction stirred at room temperature of 12 hr. The mixture was poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic  
 20 solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography, providing 252 mg (55%) of the desired methyl ester. The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (24 mg, 10%) was  
 25 obtained as a tan solid after recrystallization from MeOH. H NMR (400 MHz, DMSO) 6.89 (d, 1H), 7.28 (t, 1H), 7.37 (t, 1H), 7.61 (d, 1H), 7.81-7.86 (m, 2H), 8.01 (d, 1H), 8.11 (dd, 1H), 8.24 (t, 2H), 8.42 (d, 1H), 8.52 (t, 1H), 8.75 (d, 1H).

**5-cyano-2-([3-[(5-methoxy-1H-indol-1-yl)sulfonyl]benzoyl]amino)benzoic acid**



5-Methoxyindole (190 mg, 1.25 mmol) was dissolved in 15 ml of THF. NaH (100 mg, 60% disp. in oil, 2.5 mmol) was added and resulting suspension stirred for 1 h. Methyl 2-{{3-(chlorosulfonyl)benzoyl}amino}-5-cyanobenzoate (378 mg, 1.0 mmol) was then added and the reaction stirred at room temperature of 12 hr. The mixture was poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography, providing 236 mg (48%) of the desired methyl ester. The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (31 mg, 13%) was obtained as a white solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 3.73 (s, 3H), 6.81 (d, 1H), 6.97 (dd, 1H), 7.11 (d, 1H), 7.79 (d, 1H), 7.82 (t, 1H), 7.89 (d, 1H), 8.11 (dd, 1H), 8.21 (t, 1H), 8.42 (d, 1H), 8.48 (t, 1H), 8.76 (d, 1H).

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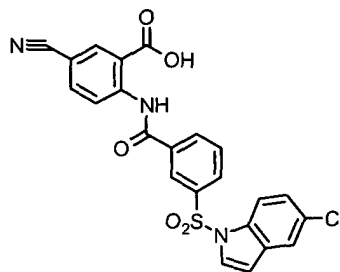
**5-cyano-2-({3-[(7-methoxy-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid** was produced using 7-Methoxyindole. <sup>1</sup>H NMR (400 MHz, DMSO) 3.84 (s, 3H), 6.79 (d, 1H), 6.83 (d, 1H), 7.29 (t, 1H), 7.61 (d, 1H), 7.75 (d, 1H), 7.80 (m, 2H), 8.17 (d, 1H), 8.28 (d, 1H), 8.32 (d, 1H), 8.56 (t, 1H), 8.73 (d, 1H).

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**5-cyano-2-({3-[(6-methoxy-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid** was produced using 6-Methoxyindole. <sup>1</sup>H NMR (300 MHz, DMSO) 3.85 (s, 3H), 6.78 (d, 1H), 6.89 (dd, 1H), 7.47-7.49 (m, 2H), 7.71 (d, 1H), 7.79-7.85 (m, 2H), 8.20 (d, 1H), 8.29 (d, 1H), 8.34 (d, 1H), 8.59 (t, 1H), 8.75 (d, 1H).

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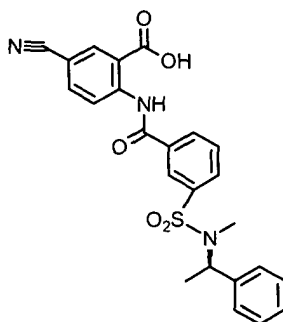
**2-({3-[(5-chloro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-cyanobenzoic acid**



5-Chloroindole (190 mg, 1.25 mmol) was dissolved in 15 ml of THF. NaH (100 mg, 60% disp. in oil, 2.5 mmol) was added and resulting suspension stirred for 1 h. Methyl 2-{{3-(chlorosulfonyl)benzoyl}amino}-5-cyanobenzoate (378 mg, 1.0 mmol) was then added and the reaction stirred at room temperature of 12 hr. The mixture was poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography, providing 311 mg (63%) of the desired methyl ester. The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (27 mg, 9%) was obtained as a white solid after recrystallization from MeOH. H NMR (400 MHz, DMSO) 6.88 (d, 1H), 7.42 (dd, 1H), 7.71 (d, 1H), 7.85 (t, 1H), 7.94 (d, 1H), 8.02 (d, 1H), 8.12 (dd, 1H), 8.25 (m, 2H), 8.43 (d, 1H), 8.52 (t, 1H), 8.76 (d, 1H).

**5-cyano-2-({3-[(5-fluoro-1H-indol-1-yl)sulfonyl]benzoyl}amino)benzoic acid** was produced utilizing 5-Fluoroindole. H NMR (400 MHz, DMSO) 6.89 (d, 1H), 7.23 (dt, 1H), 7.44 (dd, 1H), 7.85 (t, 1H), 7.96 (d, 1H), 8.00 (dd, 1H), 8.13 (dd, 1H), 8.22 (d, 1H), 8.27 (d, 1H), 8.37 (d, 1H), 8.51 (m, 2H).

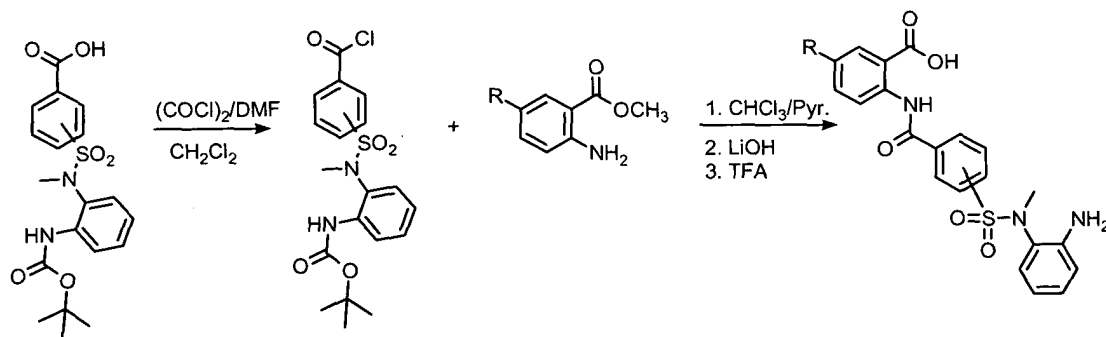
**5-cyano-2-{{3-[(1R)-1-phenylethyl]sulfonyl}benzoyl}amino}benzoic acid**



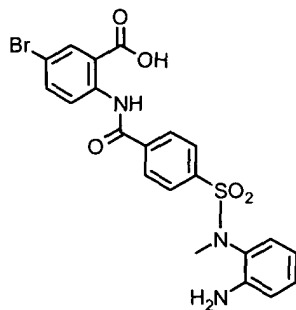
Methyl 2-{{3-(chlorosulfonyl)benzoyl}amino}-5-cyanobenzoate (378 mg, 1.0 mmol) was dissolved in 15 mL of  $\text{CHCl}_3$ . N-methyl-N-[(1R)-1-phenylethyl]amine (270 mg, 2.0 mmol) and  $\text{Et}_3\text{N}$  (1 mL) were then added and the reaction stirred at room temperature for 12 hr. The mixture was poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography, providing 400 mg (84%) of the desired methyl ester. The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over  $\text{Na}_2\text{SO}_4$  and then concentrated *in vacuo*. The title compound (300 mg, 77%) was obtained as a white solid after recrystallization from MeOH.  $^1\text{H}$  NMR (300 MHz, DMSO) 1.23 (d, 3H), 2.61 (s, 3H), 5.22 (q, 1H), 7.26-7.35 (m, 5H), 7.87 (t, 1H), 8.12 (d, 1H), 8.13 (d, 1H), 8.25 (d, 1H), 8.36 (s, 1H), 8.42 (d, 1H), 8.83 (d, 1H), 12.55 (s, 1H).

**5-cyano-2-{{3-({methyl[(1S)-1-phenylethyl]amino}sulfonyl)benzoyl}amino}benzoic acid** was produced from N-methyl-N-[(1S)-1-phenylethyl]amine.  $^1\text{H}$  NMR (300 MHz, DMSO) 1.23 (d, 3H), 2.61 (s, 3H), 5.22 (q, 1H), 7.26-7.35 (m, 5H), 7.87 (t, 1H), 8.12 (d, 1H), 8.13 (d, 1H), 8.25 (d, 1H), 8.36 (s, 1H), 8.42 (d, 1H), 8.83 (d, 1H), 12.55 (s, 1H).

Scheme 1.4

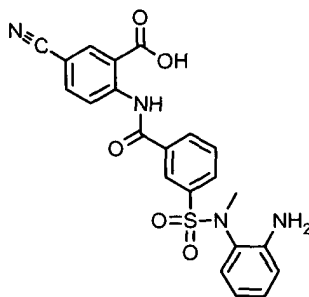


**2-[(4-{{(2-aminophenyl)(methyl)amino}sulfonyl}benzoyl)amino]-5-bromobenzoic acid**



- 4-{{2-[(tert-butoxycarbonyl)amino]phenyl}(methyl)amino)sulfonyl}benzoic (406 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs.
- 5 The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$
- 10 and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography to provide 346 mg of the desired methyl ester (56%). The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification. The resulting solid was dried in the air then dissolved in  $\text{CH}_2\text{Cl}_2$ /TFA and stirred for 10 additional hours. The solvent was removed *in vacuo* and the remaining solid was
- 15 recrystallized from MeOH to give the title compound (163 mg, 58%) as a white solid.  $^1\text{H}$  NMR (400 MHz, DMSO) 3.12 (s, 3H), 6.36-6.43 (m, 2H), 6.78 (d, 1H), 6.99-7.04 (m, 1H), 7.83-7.93 (m, 3H), 8.13-8.16 (m, 2H), 8.28-8.29 (m, 1H), 8.60 (t, 1H), 12.21 (s, 1H).

- 20 **2-[(3-{{(2-aminophenyl)(methyl)amino)sulfonyl}benzoyl}amino]-5-cyanobenzoic acid**



3-{{2-[(tert-butoxycarbonyl)amino]phenyl}(methyl)amino)sulfonyl}benzoic (406 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7

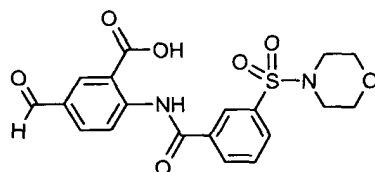
mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-cyanobenzoate (176 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography to provide 344 mg of the desired methyl ester (61%). The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification. The resulting solid was dried in the air then dissolved in  $\text{CH}_2\text{Cl}_2$ /TFA and stirred for 10 additional hours. The solvent was removed *in vacuo* and the remaining solid was recrystallized from MeOH to give the title compound (34 mg, 12%) as a white solid.  $^1\text{H}$  NMR (400 MHz, DMSO) 3.11 (s, 1H), 6.37 (m, 2H), 6.76 (d, 1H), 7.00 (m, 1H), 7.84-7.93 (m, 2H), 8.31 (dd, 1H), 8.31 (m, 2H), 8.42 (d, 1H), 8.83 (d, 1H), 12.55 (s, 1H).

#### Preparation of Methyl 2-amino-5-formylbenzoate

To a solution of methyl anthranilate (7.75 g, 51.3 mmol, Aldrich) in DMF (50 mL) was added NIS (11.5 g, 51.3 mmol, Aldrich). The solution was stirred for 63 hours before being added to a separatory funnel with 200 mL of MTBE and washed with 5 X 200 mL of water. The organics were dried over  $\text{MgSO}_4$  and evaporated yielding 13.8 g of tan solid as methyl 2-amino-5-iodobenzoate. A mixture of methyl 2-amino-5-iodobenzoate (3.13 g, 11.3 mmol) and tetrakis(triphenylphosphine)palladium(0) (282 mg, 0.244 mmol, Strem) was placed under 1 atm of CO. THF (20 mL) was added, and the solution was heated to 60 °C. Tri-*n*-butyltin hydride (3.7 mL, 12.7 mmol, Aldrich) was added dropwise with rapid stirring over 4 hours. The dark orange solution was heated a further 45 minutes and then added to a separatory funnel with 150 mL of EtOAc. This solution was washed with 2 X 150 mL of saturated aqueous  $\text{NaHCO}_3$  followed by 100 mL of brine. It was dried over  $\text{MgSO}_4$  and evaporated leaving a brown oil that was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from  $\text{CH}_2\text{Cl}_2$  to 5% EtOAc in  $\text{CH}_2\text{Cl}_2$  as eluent. This chromatography failed to remove all of the tin, so the product was re-

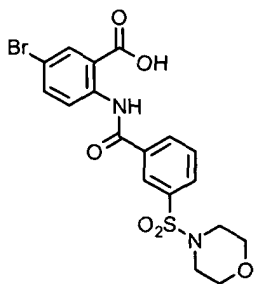
chromatographed using a Biotage Flash 40 M silica cartridge with 5% EtOAc in  $\text{CH}_2\text{Cl}_2$  as eluent. Yield was 863 mg of white solid.

**5-Formyl-2-{{3-(morpholin-4-ylsulfonyl)benzoyl}amino}benzoic acid**



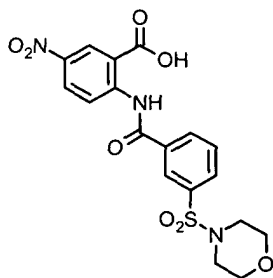
5 To 3-(morpholin-4-ylsulfonyl)benzoic acid (1.12 g, 4.13 mmol) in  $\text{CH}_2\text{Cl}_2$  (60 mL) was added DMF (20  $\mu\text{L}$ ) and oxalyl chloride (450  $\mu\text{L}$ , 5.16 mmol). The mixture was stirred for 3.75 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (20 mL), and methyl 2-amino-5-formylbenzoate (637 mg, 3.56 mmol) in pyridine (8 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of  $\text{CH}_2\text{Cl}_2$ . This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 5% EtOAc in  $\text{CH}_2\text{Cl}_2$  to 10% EtOAc in  $\text{CH}_2\text{Cl}_2$  as eluent. Yield was 636 mg of yellow solid as the methyl ester. To a mixture of the corresponding methyl ester (318 mg, 0.735 mmol) in dioxane (15 mL) was added 1 M aqueous sodium hydroxide (1.5 mL). The mixture was stirred at room temperature for 2 hours. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of EtOAc. The EtOAc was washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. It was then dried over  $\text{MgSO}_4$  and evaporated. The residue was recrystallized from hot ethanol. The solids were washed with ethanol followed by heptane and then dried at 100  $^\circ\text{C}$  under vacuum yielding 64 mg of tan solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.65 (s, 1 H), 10.00 (s, 1 H), 8.89 (d,  $J$  = 8.7 Hz, 1 H), 8.60 (d,  $J$  = 2.1 Hz, 1 H), 8.29-8.33 (m, 2 H), 8.20 (dd,  $J$  = 8.7, 2.1 Hz, 1 H), 8.03 (d,  $J$  = 8.1 Hz, 1 H), 7.93 (t,  $J$  = 7.8 Hz, 1 H), 3.63-3.68 (m, 4 H), 2.92-2.97 (m, 4 H).

**5-bromo-2-{{3-(morpholin-4-ylsulfonyl)benzoyl}amino}benzoic acid**



3-(morpholin-4-ylsulfonyl)benzoic acid (271 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-bromobenzoate (230 mg, 1.0 mmol) was added followed by pyridine (1 mL). The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (20% EtOAc in hexane) to provide 367 mg of the desired methyl ester (76%). The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over  $\text{Na}_2\text{SO}_4$  and then concentrated *in vacuo*. The title compound (328 mg, 92%, 70% overall) was obtained as a white solid after recrystallization from MeOH.  $^1\text{H}$  NMR (400 MHz, DMSO) 2.93 (m, 4H), 3.65 (m, 4H), 7.88 (dd, 1H), 7.90 (d, 1H), 8.00 (d, 1H), 8.13 (d, 1H), 8.25-8.29 (m, 2H), 8.59 (d, 1H), 12.21 (s, 1H).

## 2-[[3-(morpholin-4-ylsulfonyl)benzoyl]amino]-5-nitrobenzoic acid



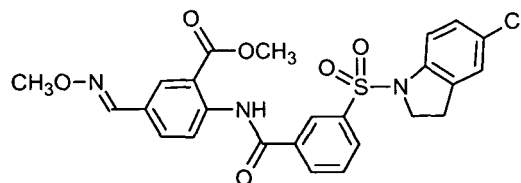
3-(morpholin-4-ylsulfonyl)benzoic acid (271 mg, 1.0 mmol) was suspended in  $\text{CH}_2\text{Cl}_2$  (10 mL) and  $(\text{COCl})_2$  added (725 mg, 5.7 mmol). A catalytic amount of DMF was then added and the mixture stirred for 4 hrs. The solvent was then removed *in vacuo* to give the acid chloride as an oil. The oil was dissolved in  $\text{CHCl}_3$  (10 mL). Methyl 2-amino-5-nitrobenzoate (196 mg, 1.0 mmol) was added followed by pyridine (1 mL).



The solution was stirred at room temperature for an additional 12 hrs then poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography (20% EtOAc in hexane) to provide 108 mg of the desired methyl ester (24%). The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (70 mg, 67%, 16% overall) was obtained as a yellow solid after recrystallization from MeOH. <sup>1</sup>H NMR (400 MHz, DMSO) 2.94 (m, 4H), 3.65 (m, 4H), 7.94 (t, 1H), 8.04 (d, 1H), 8.29-8.33 (m, 2H), 8.55 (dd, 1H), 8.80 (d, 1H), 8.91 (d, 1H), 12.76 (s, 1H)

**Methyl 2-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-formylbenzoate** was prepared as described above using 3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoic acid. **2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-formylbenzoic acid** was prepared by hydrolyzing the corresponding methyl ester. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.69 (s, 1 H), 10.00 (s, 1 H), 8.87 (d, *J* = 8.7 Hz, 1 H), 8.61 (d, *J* = 2.1 Hz, 1 H), 8.40 (s, 1 H), 8.27 (d, *J* = 7.9 Hz, 1 H), 8.19 (dd, *J* = 8.7, 2.1 Hz, 1 H), 8.09 (d, *J* = 8.5 Hz, 1 H), 7.84 (t, *J* = 7.8 Hz, 1 H), 7.53 (d, *J* = 8.5 Hz, 1 H), 7.24-7.29 (m, 2 H), 4.02 (t, *J* = 8.5 Hz, 2 H), 2.95 (t, *J* = 8.4 Hz, 2 H).

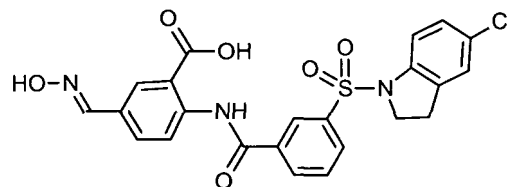
**2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-[(E)-(methoxyimino)methyl]benzoic acid**



A slurry of methyl 2-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-formylbenzoate (475 mg, 0.952 mmol) and O-methylhydroxylamine hydrochloride (526 mg, 6.30 mmol, Aldrich) in 1:1 ethanol/pyridine (25 mL) was stirred for 2 days. The mixture was then added to a separatory funnel with 120 mL of CH<sub>2</sub>Cl<sub>2</sub>. This solution was washed with 2 X 100 mL of 1 M aqueous HCl followed by 100 mL of brine. The CH<sub>2</sub>Cl<sub>2</sub> was evaporated in

the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from CH<sub>2</sub>Cl<sub>2</sub> to 2% EtOAc in CH<sub>2</sub>Cl<sub>2</sub> as eluent. Yield was 411 mg of white solid as the methyl ester. To a mixture of the corresponding methyl ester (288 mg, 0.545 mmol) in dioxane (20 mL) was added 1 M aqueous sodium hydroxide (1.5 mL). The mixture was stirred at room temperature for 4.5 hours and then in a 50 °C oil bath for 30 minutes. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of EtOAc. The EtOAc was washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. It was then dried over MgSO<sub>4</sub> and evaporated. The residue was recrystallized from hot ethanol/THF. The solids were washed with ethanol followed by heptane and then dried at 100 °C under vacuum yielding 127 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.40 (s, 1 H), 8.70 (d, *J* = 8.7 Hz, 1 H), 8.38 (s, 1 H), 8.31 (d, *J* = 2.1 Hz, 1 H), 8.30 (s, 1 H), 8.25 (d, *J* = 7.9 Hz, 1 H), 8.07 (d, *J* = 8.1 Hz, 1 H), 7.91 (dd, *J* = 8.7, 2.1 Hz, 1 H), 7.83 (t, *J* = 7.9 Hz, 1 H), 7.52 (d, *J* = 8.5 Hz, 1 H), 7.24-7.29 (m, 2 H), 4.02 (t, *J* = 8.5 Hz, 2 H), 3.91 (s, 3 H), 2.95 (t, *J* = 8.4 Hz, 2 H).

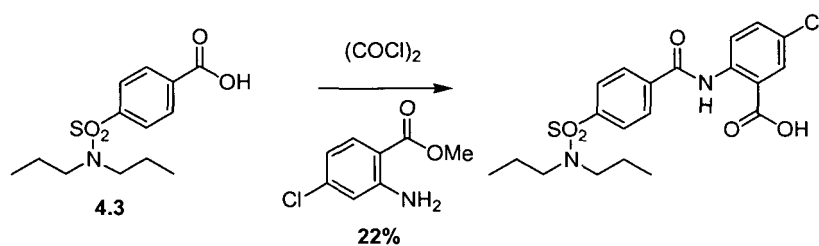
**2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-[(E)-(hydroxyimino)methyl]benzoic acid**



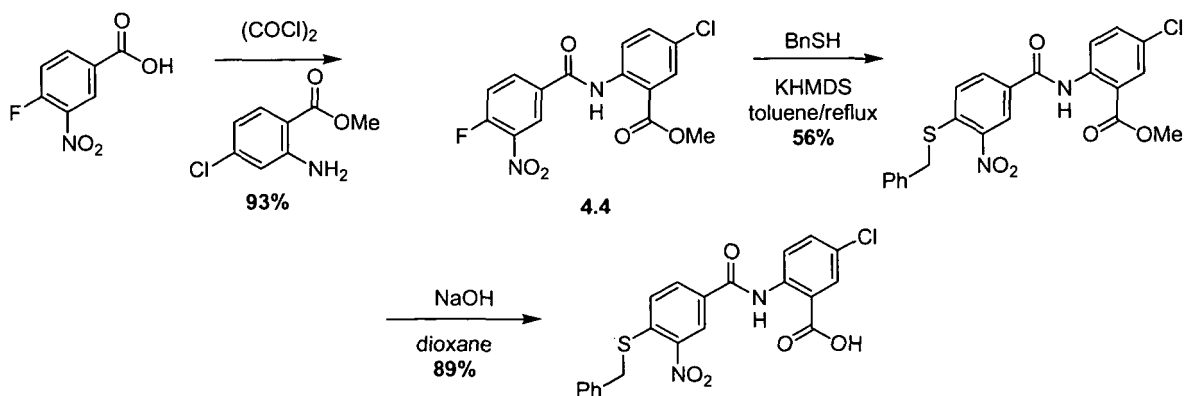
A slurry of methyl 2-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-formylbenzoate (627 mg, 1.26 mmol) and hydroxylamine hydrochloride (656 mg, 9.44 mmol, Mallinckrodt) in 1:1 ethanol/pyridine (25 mL) was stirred for 2 days. The mixture was then added to a separatory funnel with 120 mL of CH<sub>2</sub>Cl<sub>2</sub>. This solution was washed with 2 X 100 mL of 1 M aqueous HCl followed by 100 mL of brine. The CH<sub>2</sub>Cl<sub>2</sub> was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with 5% EtOAc in CH<sub>2</sub>Cl<sub>2</sub> as eluent. Yield was 478 mg of white solid as the methyl ester. To a mixture of the corresponding methyl ester (363 mg, 0.706 mmol) in dioxane (20 mL) was added 1 M aqueous sodium

hydroxide (1.5 mL). The mixture was stirred at room temperature for 4.5 hours. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of EtOAc. The EtOAc was washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. It was then dried over MgSO<sub>4</sub> and evaporated. The residue was recrystallized from hot ethanol/THF. The solids were washed with ethanol followed by heptane and then dried at 100 °C under vacuum yielding 280 mg of white solid. Because NMR and CHN analysis were consistent with this material containing residual solvent, 200 mg of the material was heated in 50 mL of methanol. Solvent was removed, and the residue was again dried at 100 °C under vacuum yielding 183 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.37 (s, 1 H), 11.31 (s, 1 H), 8.68 (d, *J* = 8.7 Hz, 1 H), 8.38 (s, 1 H), 8.29 (d, *J* = 1.9 Hz, 1 H), 8.25 (d, *J* = 7.9 Hz, 1 H), 8.20 (s, 1 H), 8.07 (d, *J* = 8.1 Hz, 1 H), 7.90 (dd, *J* = 8.8, 2.0 Hz, 1 H), 7.83 (t, *J* = 7.9 Hz, 1 H), 7.53 (d, *J* = 8.5 Hz, 1 H), 7.24-7.29 (m, 2 H), 4.01 (t, 8.5, 2 H), 2.95 (t, *J* = 8.4 Hz, 2 H).

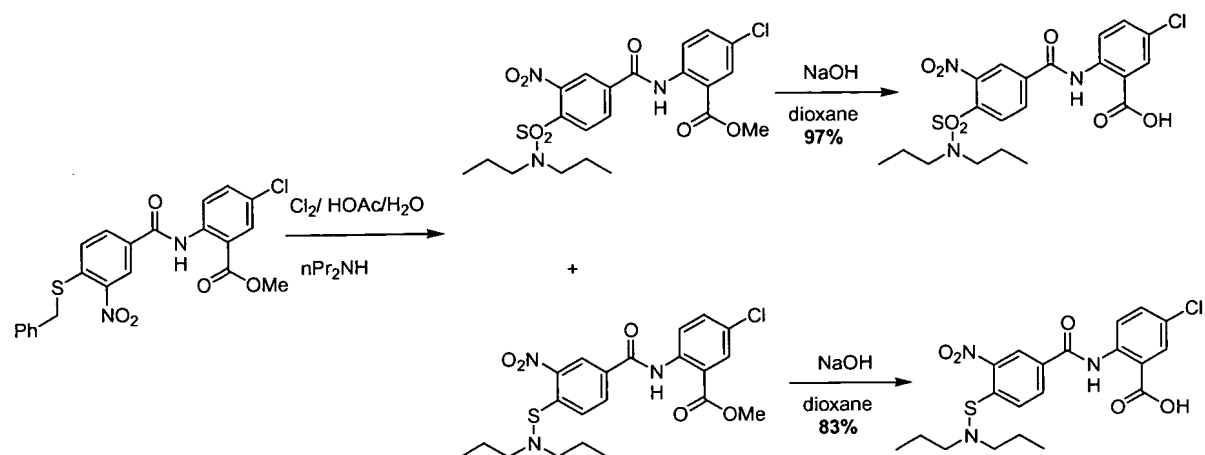
### Scheme 1.5



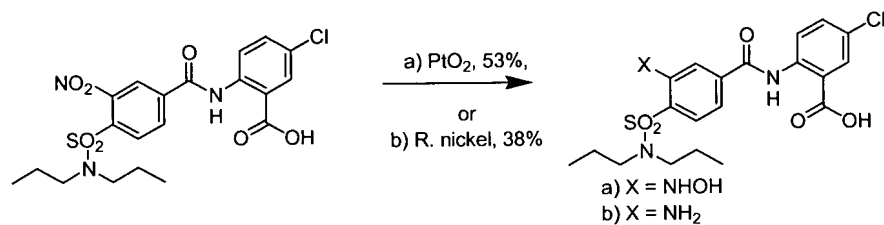
### Scheme 1.6



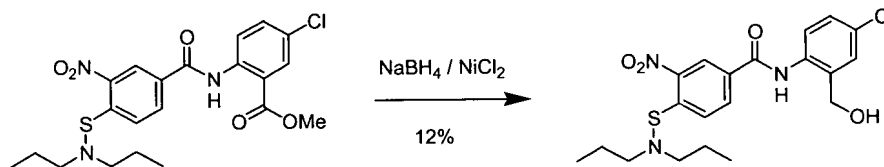
**Scheme 1.7**



**5 Scheme 1.8**

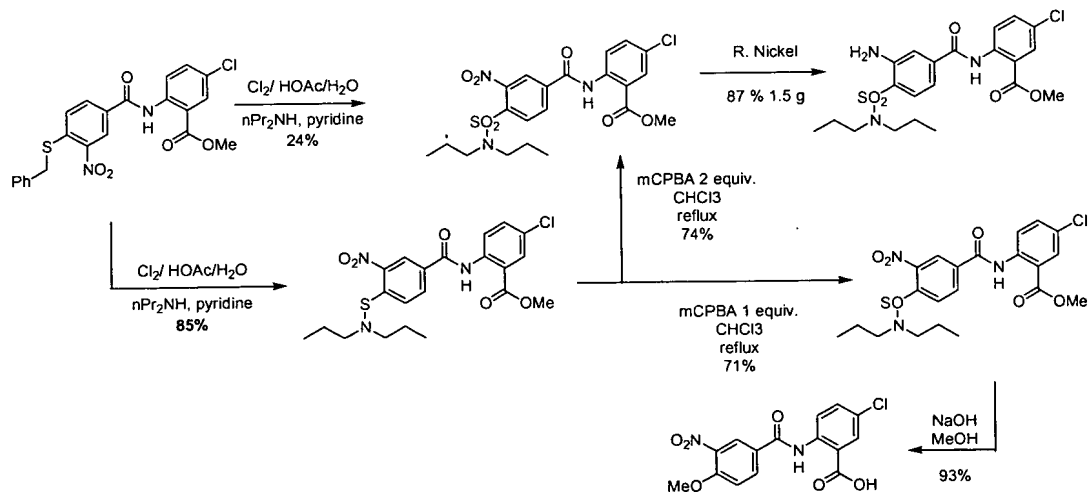


**Scheme 1.9**

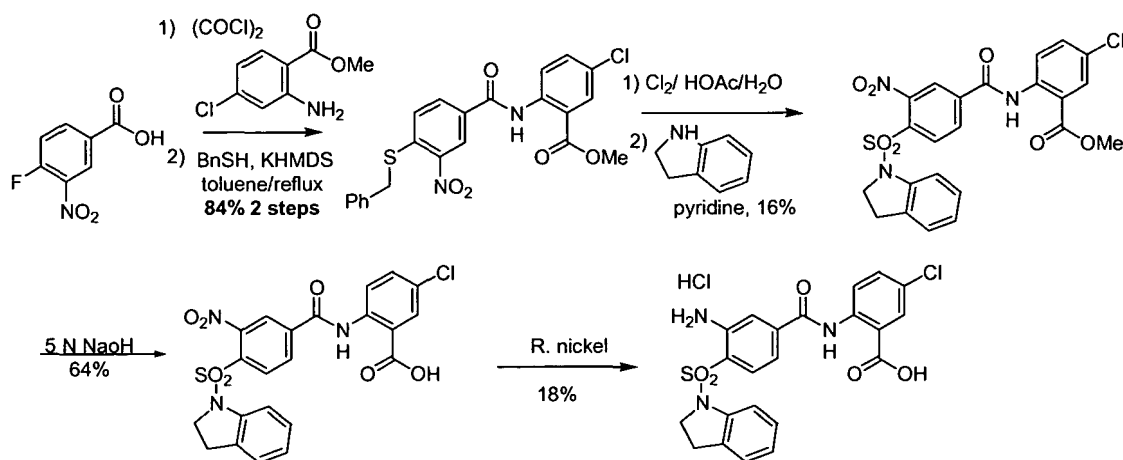


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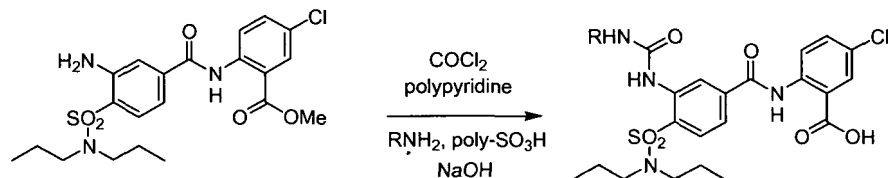
**Scheme 1.10**



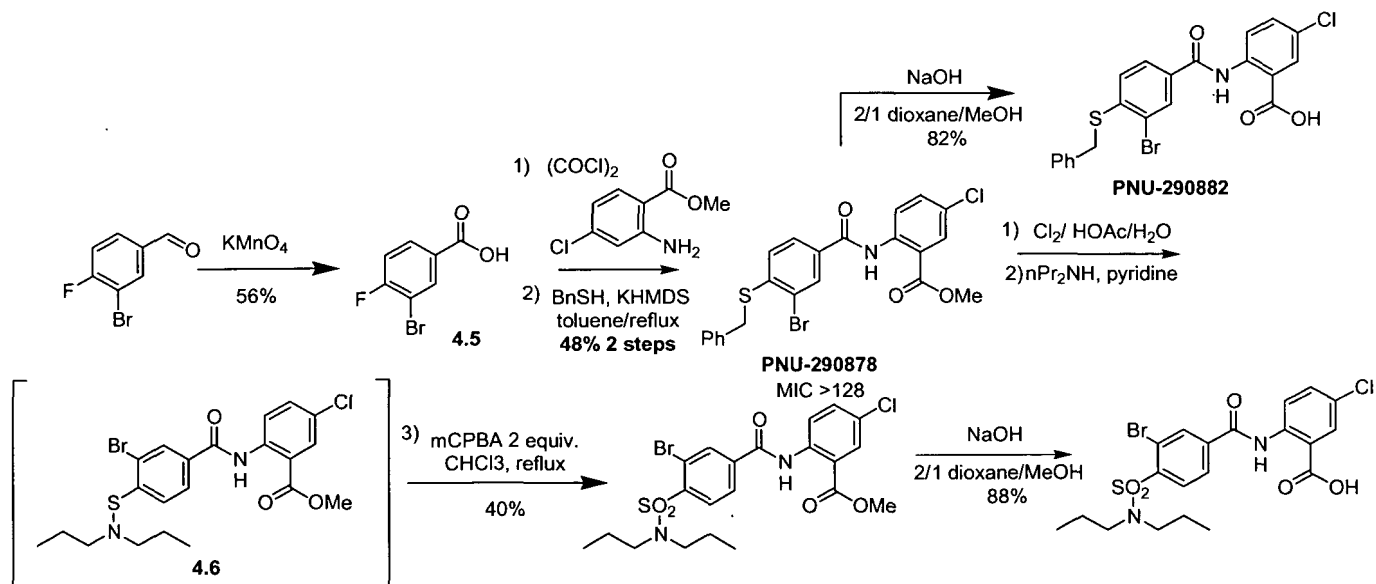
### Scheme 1.11



### Scheme 1.12



### Scheme 1.13



Compounds produced via the above-described synthetic schemes include, but are not limited to, the following:

5-Chloro-2-({4-[(dipropylamino)sulfonyl]benzoyl}amino)benzoic acid

5-Chloro-2-({4-[(dipropylamino)sulfonyl]-3-nitrobenzoyl}amino)benzoic acid

5-Chloro-2-{{4-[(dipropylamino)sulfonyl]-3-(hydroxyamino)benzoyl}amino} benzoic acid hydrochloride

2-({3-Amino-4-[(dipropylamino)sulfonyl]benzoyl}amino)-5-chlorobenzoic acid  
5 hydrochloride

2-{{4-(Benzylsulfanyl)-3-nitrobenzoyl}amino}-5-chlorobenzoic acid

5-Chloro-2-({4-[(dipropylamino)sulfanyl]-3-nitrobenzoyl}amino)benzoic acid

Methyl 5-chloro-2-({4-[(dipropylamino)sulfinyl]-3-nitrobenzoyl}amino)benzoate

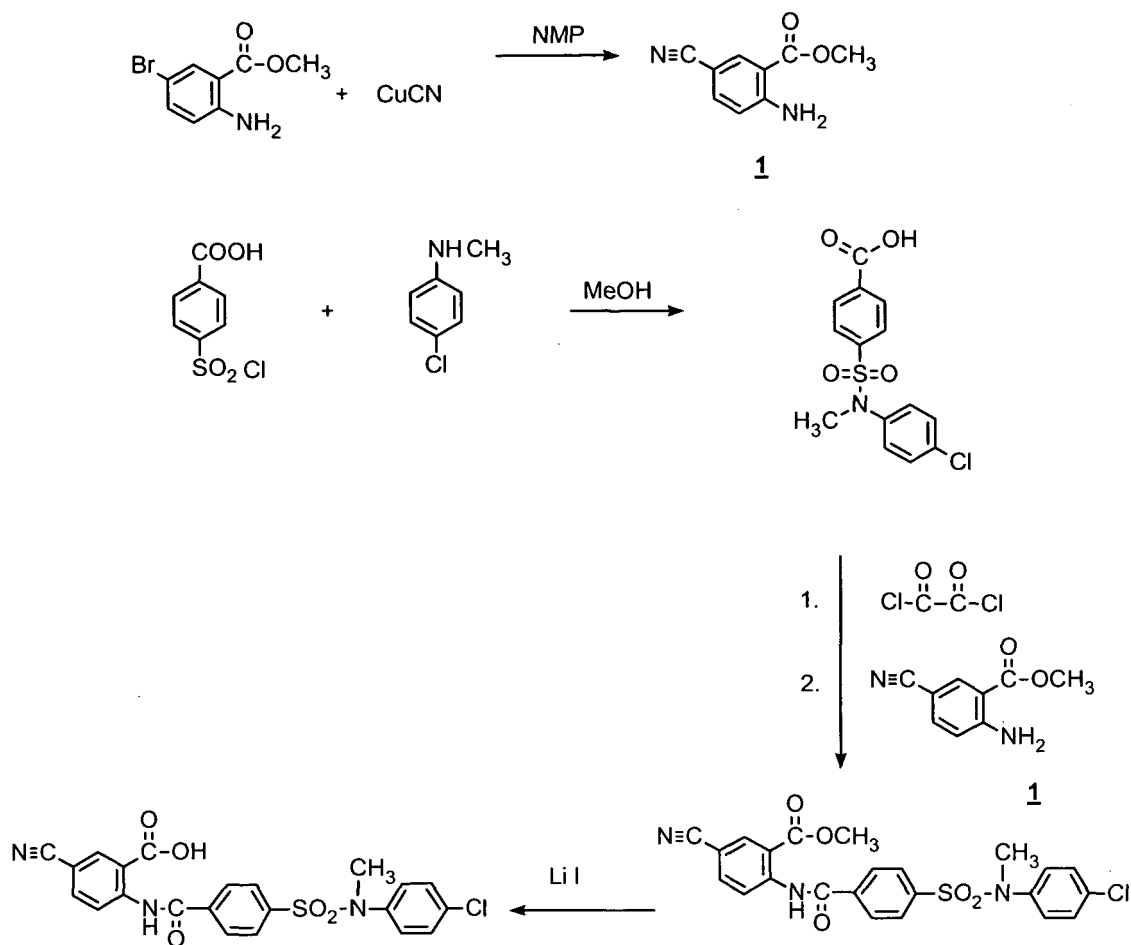
10 5-Chloro-2-{{4-(2,3-dihydro-1H-indol-1-ylsulfonyl)-3-nitrobenzoyl}amino} benzoic acid

Cyano 2-{{3-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl}amino}-5-ethynylbenzoic acid

Methyl 2-({3-amino-4-[(dipropylamino)sulfonyl]benzoyl}amino)-5-chlorobenzoate

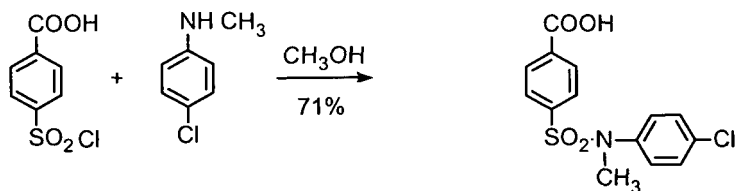
15 2-({3-Bromo-4-[(dipropylamino)sulfonyl]benzoyl}amino)-5-chlorobenzoic acid

### Scheme 1.16



### Preparation of 4-([4-chloro(methyl)anilino]sulfonyl)benzoic acid

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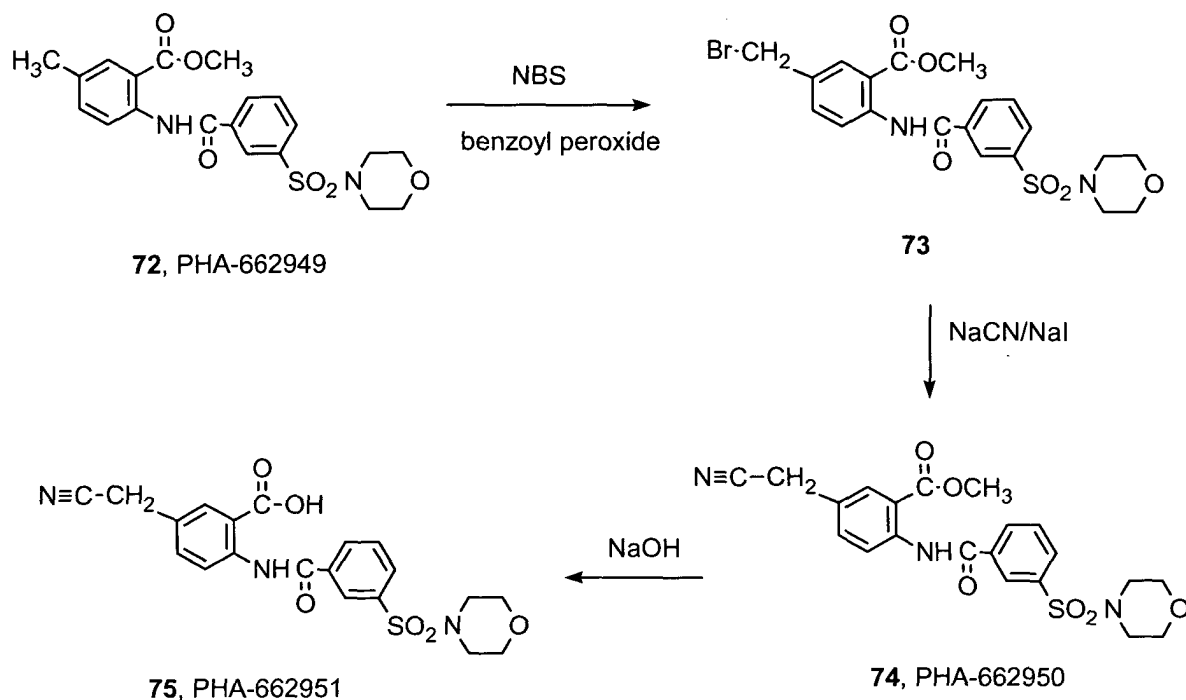


10

A solution of 4-chloro-N-methylaniline (10.0 g, 0.0706 mol, 1.1 eq) and triethylamine (7.78 g, 0.0770 mol, 1.2 eq) in 140 mL of methanol, cooled in an ice bath at 0-5°C, was treated portionwise over a one minute period with solid 4-chlorosulfonyl benzoic acid (14.2 g, 0.0642 mol, 1.0eq). After the addition was complete, the cooling bath was removed and the reaction mixture was stirred under a nitrogen atmosphere while

warming to room temperature on its own. After 5.5 h, the contents were poured into 270 mL of ice water containing 130 mL of 3 N NaOH, washed the milky solution with methylene chloride (2 X 100 mL), acidified the aqueous layer with 35 mL of concentrated HCl. After cooling the mixture in an ice bath, the white precipitated product was collected and dried in a vacuum oven at 70°C overnight to yield 14.92 g (71%) of **2**. <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 13.53 (brs, 1 H), 8.11 (dd, *J* = 2, 7 Hz, 2 H), 7.63 (dd, *J* = 2, 7 Hz, 2 H), 7.42 (dd, *J* = 2, 7 Hz, 2 H), 7.14 (dd, *J* = 2, 7 Hz, 2 H), 3.15 (s, 3 H) ppm.

**Scheme 1.17**



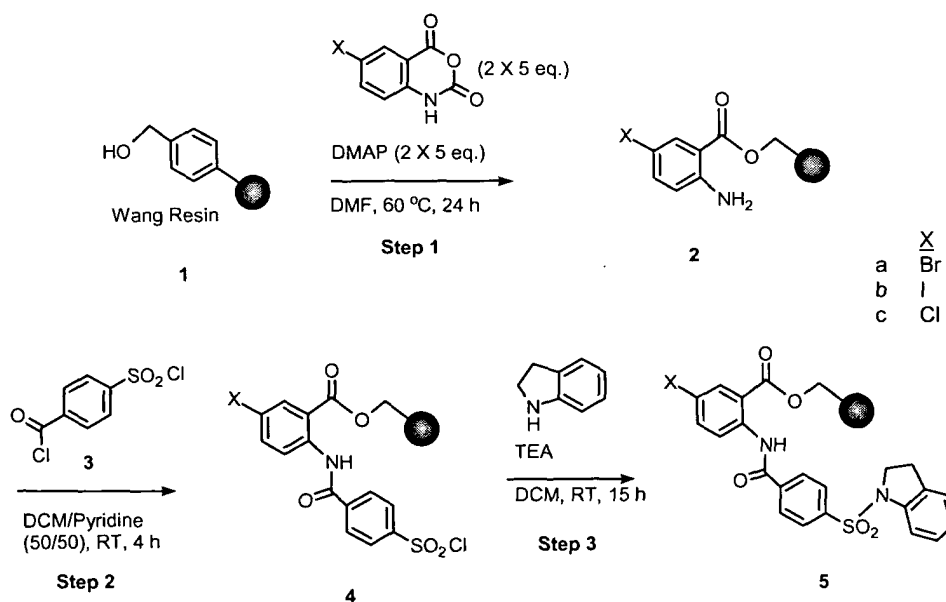
To 21 mL of carbon tetrachloride at room temperature was added benzoyl peroxide (0.095 g, 0.393 mmol, 0.10 eq). The solution was slowly heated to reflux at which time N-bromosuccinimide (0.769 g, 4.32 mmol, 1.1 eq) was added at once followed by a slurry of compound **72** (1.64 g, 3.93 mmol, 1.0 eq) in 9 mL of carbon tetrachloride plus 6 mL of carbon tetrachloride as a rinse. Vigorous refluxing was continued for 2 h, the reaction mixture filtered hot and the solids rinsed with additional hot carbon tetrachloride. The filtrate was concentrated at reduced pressure to give more than theoretical amount of crude bromomethyl compound **73**. This was dissolved in 35 mL of acetone, treated with NaCN (0.289 g, 5.90 mmol, 1.5 eq) and



NaI (0.029 g, 0.197 mmol, 0.05 eq) and the mixture refluxed for 24 h. An additional 0.50 eq (0.096 g) of NaCN was added and refluxing continued for 3 h longer. The cooled reaction mixture was filtered, the filtrate concentrated at reduced pressure, the residue dissolved in ethyl acetate and washed successively with 10 mL of water and 10 mL of 50% saturated brine. The combined aqueous washings were back extracted once with ethyl acetate, the combined organic extracts dried with anhydrous sodium sulfate and the filtrate concentrated *in vacuo*. Chromatography with 100 g of silica gel, packed and eluted with acetone-methylene chloride-heptane (1:4:5), afforded cyanomethyl ester **74** in 20% yield (based on **72**) as a white solid. Base hydrolysis of **74** (0.297 g, 0.670 mmol) in 4 mL of methylene chloride, 4 mL of methanol and 1 mL of water using 1N NaOH (3.02 mL, 4.5 eq) at room temperature gave a 55% yield of acid **75** as a white solid. **73**: TLC (silica gel GF):  $R_f = 0.36$  acetone-methylene chloride-hexane(1:3:6);  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  8.89 (d,  $J = 7$  Hz, 1 H), 8.41 (t,  $J = 1$  Hz, 1 H), 8.27 (m, 1 H), 8.14 (d,  $J = 2$  Hz, 1 H), 7.97 (m, 1 H), 7.75 (t,  $J = 6$  Hz, 1 H), 7.66 (dd,  $J = 2, 6$  Hz, 1 H), 4.52 (s, 2 H), 3.98 (s, 3 H), 3.78 (t,  $J = 3$  Hz, 4 H), 3.10 (t,  $J = 4$  Hz, 4 H) ppm.

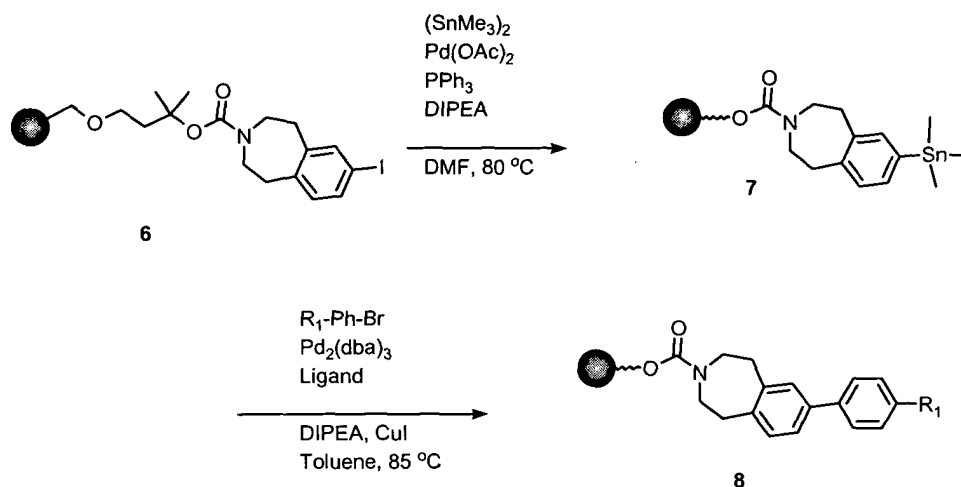
Scheme 1.18 outlines the solid phase synthesis of halogenated anthranilic acid substrates **5**.

**Scheme 1.18**



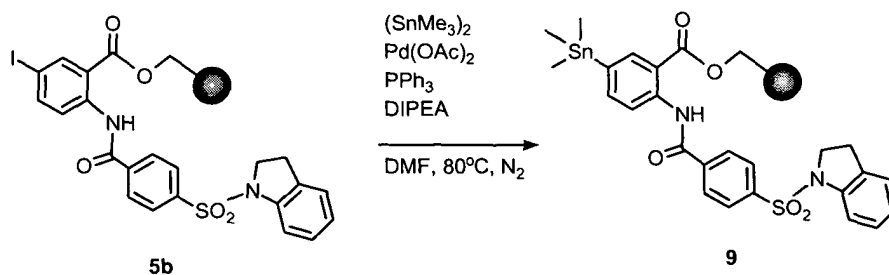
Resin bound iodide **6** was stannylated using the conditions shown in Scheme 10.2. Hunigs base, although not directly involved in the reactions, was used as a proton scavenger. A library based on this template was successfully prepared using Suzuki cross-coupling conditions.

**Scheme 1.19**



Applying the Stille conditions to the template, stannylated product **9** was prepared from iodide **5b**. The reaction was monitored via observance of the protodestannylation product after TFA cleavage from resin. Stannylation of the corresponding solid-phase bromide **5a** was less successful.

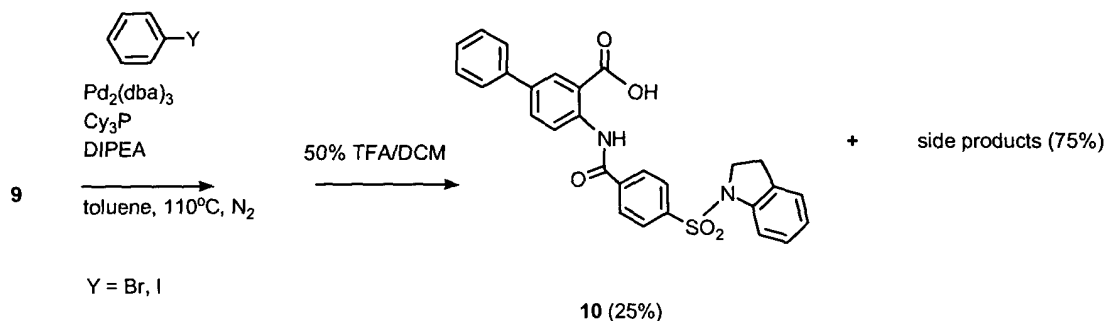
**Scheme 1.20**



Attempts at coupling aryl bromides and iodides with the stannylated resin gave some product, but not in quantities suitable for library production (Scheme 1.21). Protodestannylation and homocoupling were the major competing reactions, leaving

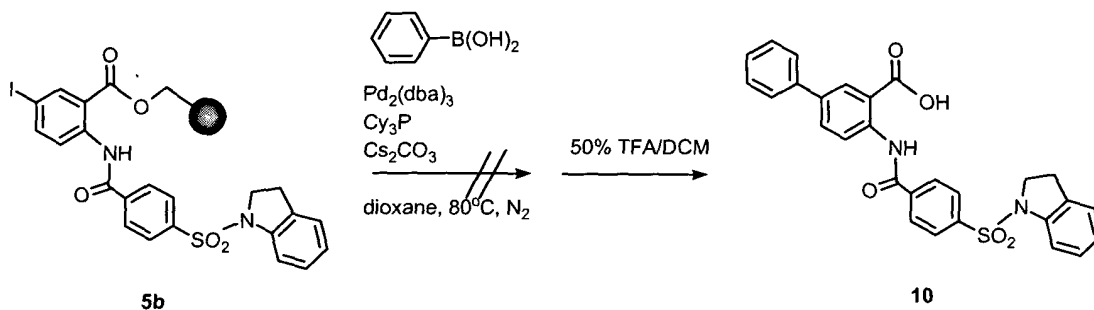
product purities in the 25 % range. The reactions were monitored by HPLC (at 210 nm), and product identities were confirmed by LC/MS.

**Scheme 1.21**



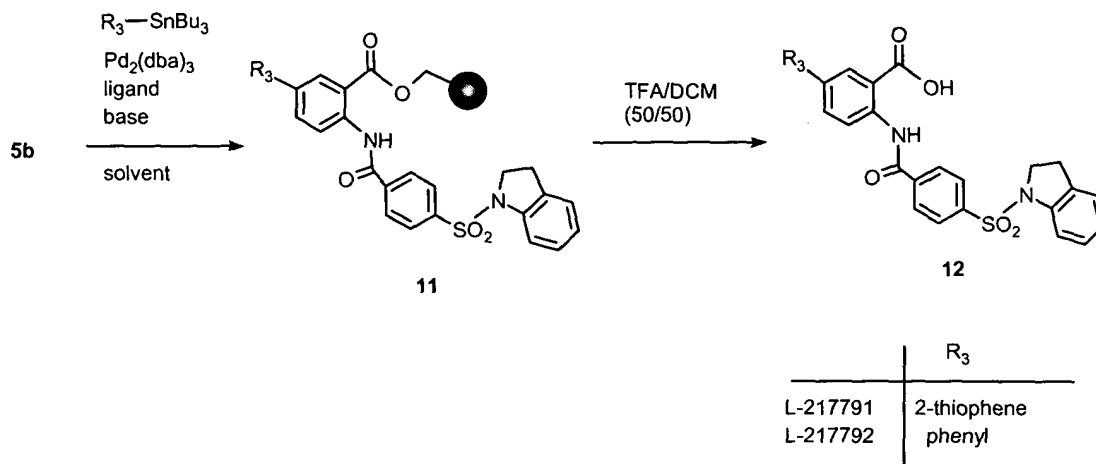
Suzuki coupling chemistry was conducted under the conditions shown in Scheme 1.22.

**Scheme 1.22**



The cross-coupling reaction from the other direction is shown in Scheme 1.23, in which purchased aryl tin compounds were coupled with the resin-bound iodide.

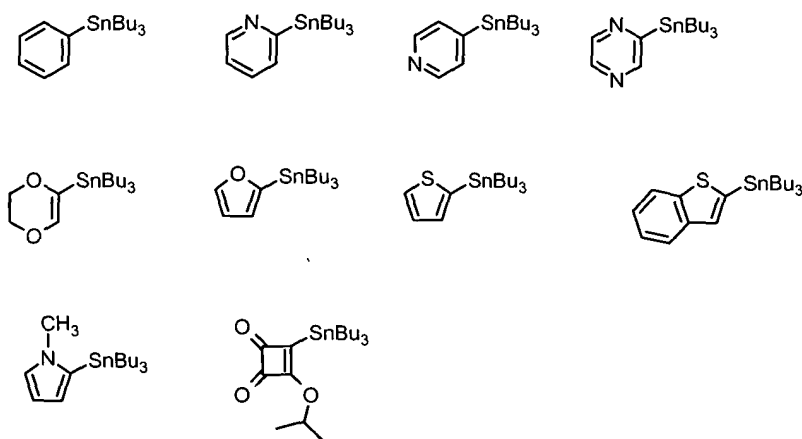
**Scheme 1.23**



The best results in the case of tributylphenyl tin were obtained in toluene with 1,1'-bis(diphenylphosphino)-ferrocene as ligand and a reaction time of 2.5 hours at 115 °C.

In the case of 2-(tributylstannyl) thiophene, toluene was the solvent of choice and tricyclohexylphosphine, triphenyl arsine, and 1,1'-bis(diphenylphosphino)-ferrocene worked equally well after 2.5 hours at 115 °C.

**Table 1.1:** Commercially Available Aryl Tin Compounds

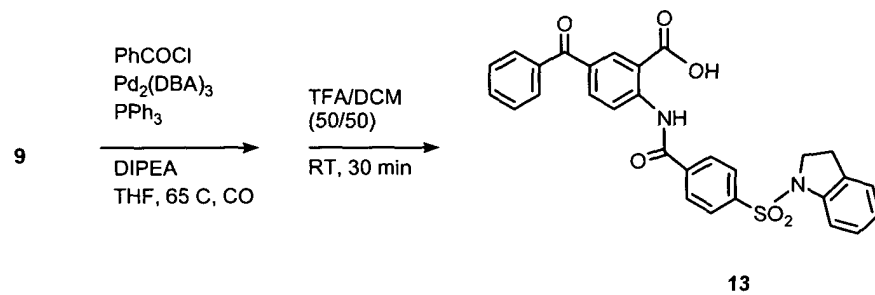


### Installation of Ketones via Palladium-Catalyzed Coupling with Acid Chlorides

Acid chlorides were coupled with **9** (see scheme 1.20) using similar, but milder conditions (Scheme 10.7). The ketone product (**13**) was produced using triphenylphosphine as ligand and THF as solvent in 75 % yield and 70 % purity. A

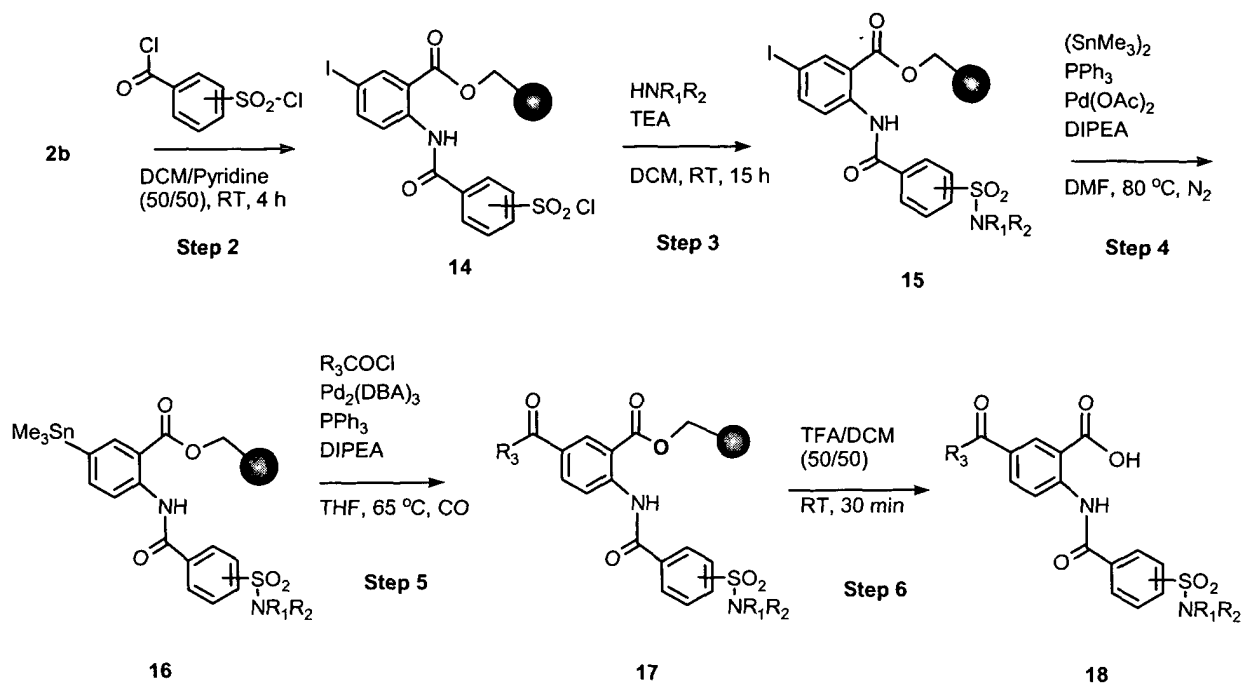
carbon monoxide atmosphere was used to eliminate small amounts of the corresponding aryl-aryl product formation (**12**), while Hunigs base was employed as the proton scavenger to help avoid protodestannylation.

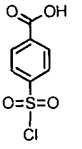
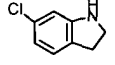
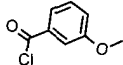
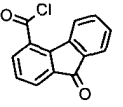
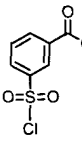
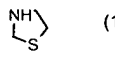
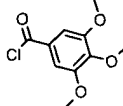
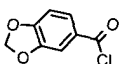
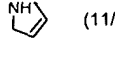
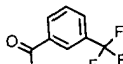
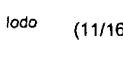
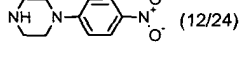
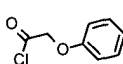
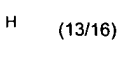
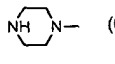
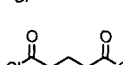
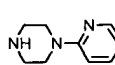
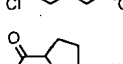


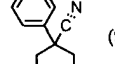
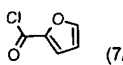
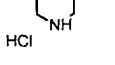
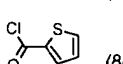
**Scheme 1.24**



5

## Scheme 1.25

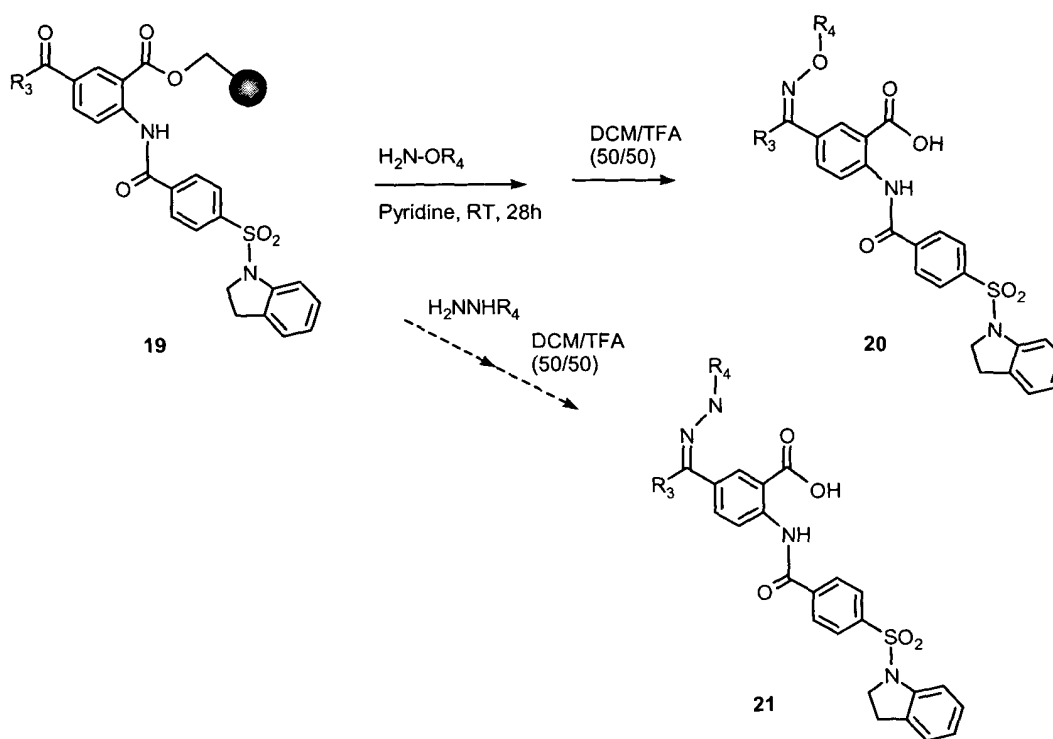
5 **Table 1.2:** Diversity Elements (no. of  $\geq 70$  % pure products/no. attempted)

sulfonyl chlorides	amines	acid chlorides	
 (39/96)	 (12/24)	 (7/16)	 (0/16)
 (33/96)	 (12/24)	 (6/16)	 (6/16)
	 (11/24)	 (1/16)	 (11/16)
	 (12/24)	 (4/16)	 (13/16)
	 (6/24)	 (4/16)	
	 (6/24)	 (6/16)	
	 (9/24)	 (3/16)	
	 (4/24)	 (7/16)	
	 (4/24)	 (8/16)	

**Derivatization of Aryl Ketones: Derivatizing the ketones as oximes, alkoxyamines, hydrazones, and amines**

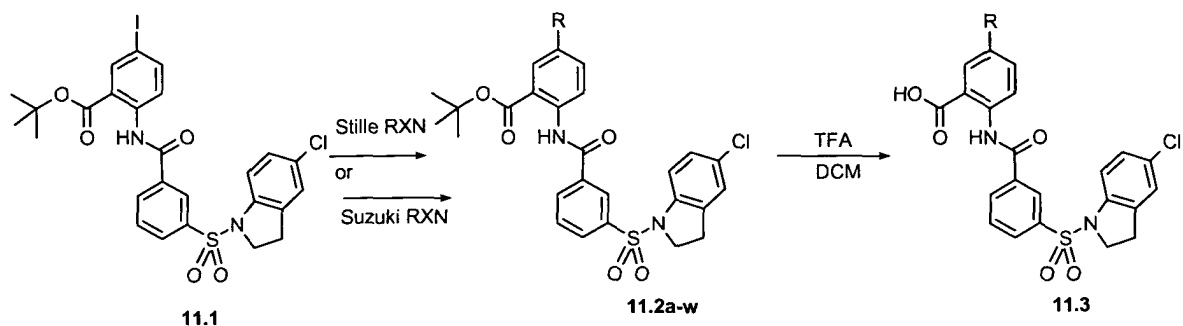
- 5 Oximes and alkoxyamines (**20**) were prepared in reasonable purities from their corresponding hydroxylamine hydrochlorides and resin **19** in pyridine (Scheme 1.26). Hydrazone, sulfonylhydrazone, and acyl-hydrazone formations (**21**) using literature conditions, however, were sluggish and could never be pushed to completion.

10 **Scheme 1.26**

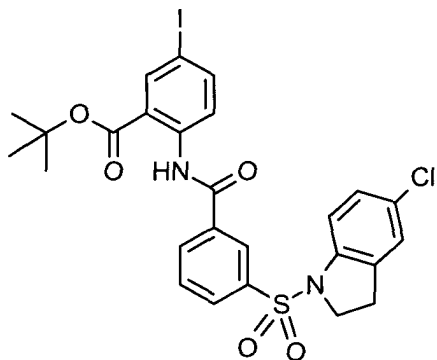


- Amines **22** were prepared on solid-phase using reductive amination. Imine formation, mediated by titanium isopropoxide, typically took four to six hours to go to completion. The sodium triacetoxy borohydride reduction was allowed to proceed overnight to give good quality amine products.
- 15

**Scheme 1.30**



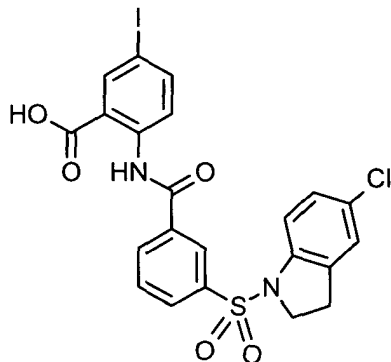
***t*-Butyl 2-((3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl)amino)-5-iodobenzoate, a, Compound 11.1**



3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoic acid (2.3 g, 6.9 mmol, 1 equivalent) and oxyl chloride (2.6 g, 20.5 mmol, 3 equivalent) were dissolved in methylene chloride (30 ml), followed by the addition of DMF (0.4 ml). Gas evolution was observed. The mixture was stirred at room temperature for 2 h later, then heptane (30 ml) was added. The solution was concentrated to dryness, and the residue was re-dissolved in DCM (30 ml), followed by the dropwise addition of PHA-561052 (2.2 g, 6.9 mmol, 1 equivalent) in DCM (20 ml) and pyridine (1.2 ml). The resulting solution was stirred overnight, then diluted with MTBE (200 ml) and washed with 0.1N HCl, 1N NaOH, brine, dried (MgSO<sub>4</sub>), filtered, and concentrated *in vacuo*. The residue was recrystallized from heptane to afford 2.4 g (55%) of 1 as a yellow solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.34 (s, 1 H), 8.67 (d, *J* = 9.0 Hz, 1 H), 8.54 (s, 1 H), 8.33 (m, 1 H), 8.24 (d, *J* = 8.5 Hz, 1 H), 7.97 (d, *J* = 8.4 Hz, 1 H), 7.88 (d, *J* = 8.5 Hz, 1 H), 7.64 (m, 2 H), 7.17 (d, *J* = 8.5 Hz, 1 H), 7.06 (s, 1 H), 4.08 (t, *J* = 8.5 Hz, 2 H), 2.95 (t, *J* = 8.4 Hz, 2 H), 1.67 (s, 9 H).



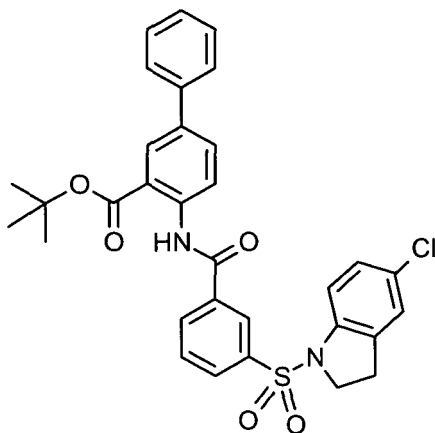
**2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-iodobenzoic acid**



5 General method E: (Hydrolysis of the alkyl ester)

Ester **11.1** (150mg, 0.24mmol) was dissolved in DCM (6 ml), followed by the addition of TFA (1.2 ml). The solution was shaken overnight, then diluted with DCM (5 ml) and heptane (1 ml). The solution was concentrated *in vacuo* to dryness, the  
 10 residue was pumped for about 1h, then triturated with methanol, filtered to afford 102mg(75%) of a white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.25 (s, 1 H), 8.44 (d, *J* = 9 Hz, 1 H), 8.33 (s, 2 H), 8.31 (m, 1 H), 8.05 (m, 2 H), 7.81 (t, *J* = 8.5 Hz, 1 H), 7.71 (d, *J* = 9 Hz, 1 H), 7.24 (m, 2 H), 4.01 (t, *J* = 8.1 Hz, 2 H), 2.95 (t, 2 H).

***t*-Butyl 4-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)[1,1'-biphenyl]-3-carboxylate, 2a**



5

**General method F:**

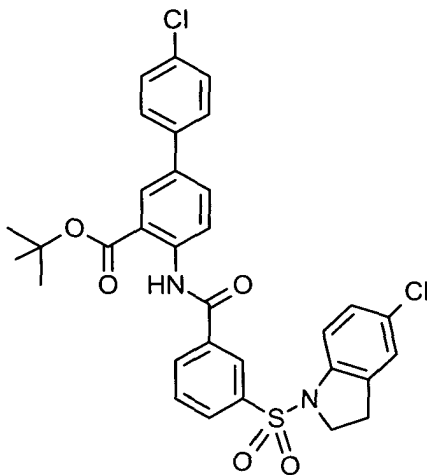
Ester **11.1** (150 mg, 0.235 mmol) and tetrakis(triphenylphosphine) palladium(0) (13.6 mg, 0.01175 mmol) were placed in a 50ml one-necked round bottom flask. The system was evacuated and filled with argon several times. Then tributylstannylbenzene (91.75 mg, 0.25 mmol) in toluene (10 ml) was added. The resulting solution was heated at 100°C overnight, cooled to room temperature, then KF (87mg, ) was added. The mixture was stirred at room temperature for 2h, filtered through celite. The filtrate was concentrated *in vacuo* and the residue was purified by silica gel chromatography (EtOAc/heptane 1/25, 1/10) to afford 120 mg (88%) of **11.2a** as a yellow solid.

15

**General method G:**

Ester **11.1** (150 mg, 0.235 mmol) and dichlorobis(triphenylphosphine) palladium (II) (8.4 mg, 0.012 mmol) were placed in a 50 ml one-necked round bottom flask. The system was evacuated and filled with argon several times. Then tributylstannylbenzene (91.7 mg, 0.25 mmol) in THF (10 ml) was added. The resulting solution was heated at 80°C overnight, cooled to room temperature, KF (87 mg) was added. The mixture was stirred at room temperature for 2h, filtered through celite. The filtrate was concentrated *in vacuo* and the residue was purified by silica gel chromatography (EtOAc/heptane 1/25, 1/10) to afford 101 mg (74%) of **11.2a** as a yellow solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 11.60 (s, 1 H), 8.42 (s, 1 H), 8.35 (d, *J* = 9 Hz, 1 H), 8.27 (d, *J* = 8 Hz, 1 H), 8.15 (d, *J* = 2 Hz, 1 H), 8.06 (d, *J* = 8 Hz, 1 H), 7.98 (d, *J* = 9 Hz, 1 H), 7.84 (t, *J* = 8 Hz, 1 H), 7.70 (d, *J* = 7 Hz, 2 H), 7.50 (m, 3 H), 7.41 (t, *J* = 7 Hz, 1 H), 7.25 (d, *J* = 7 Hz, 2 H), 4.05 (t, *J* = 8 Hz, 2 H), 2.97 (t, *J* = 8 Hz, 2 H), 1.53 (s, 9 H).

***t*-Butyl 4'-chloro-4-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)[1,1'-biphenyl]-3-carboxylate, **11.2c****

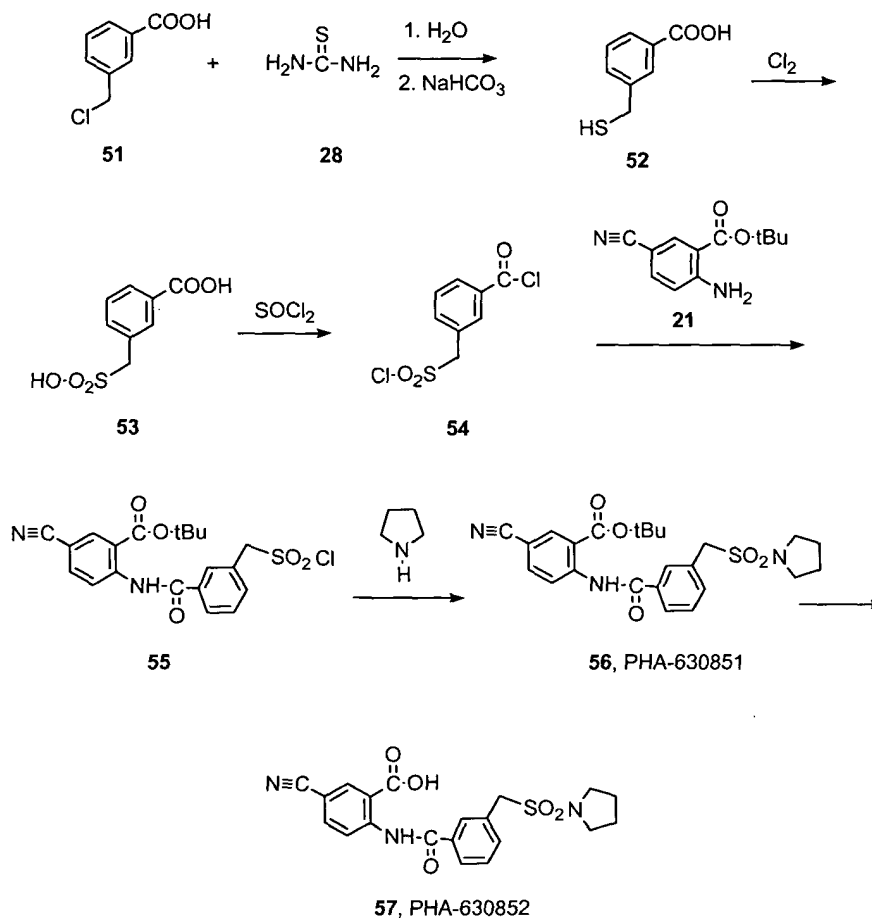


General method H:

Ester **11.1** (160 mg, 0.25 mmol), tetrakis(triphenylphosphine) palladium(0) (14.5 mg, 0.0125 mmol), sodium carbonate (101 mg, 0.95 mmol) and 4-chlorobenzenboronic acid (43 mg, 0.275 mmol) were placed in a 100ml one-necked round bottom flask. The system was evacuated and filled with argon several times. Then THF (50 ml) and distilled water (5 ml) were added. The solution was heated at reflux temperature for 20h, the solvent was removed in *vacuo* and residue was purified by silica gel

chromatography (EtOAc/heptane 1/25, 1/10) to get 92 mg (59%) of **11.2c** as a yellow solid.  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ )  $\delta$  12.40 (s, 1 H), 8.93 (d,  $J = 9$  Hz, 1 H), 8.59 (s, 1 H), 8.28 (d,  $J = 8$  Hz, 1 H), 8.24 (d,  $J = 2.3$  Hz, 1 H), 7.95 (d,  $J = 8$  Hz, 1 H), 7.80 (dd,  $J = 2.5, 8.2$  Hz, 1 H), 7.64 (m, 6 H), 7.20 (d,  $J = 8$  Hz, 1 H), 7.08 (s, 1 H), 4.12 (t,  $J =$   
 5 8 Hz, 2 H), 2.98 (t,  $J = 8$  Hz, 2 H), 1.71 (s, 9 H).

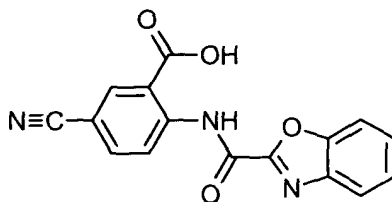
**5-cyano-2-({3-[(1-pyrrolidinylsulfonyl)methyl]benzoyl}amino)benzoic acid PHA-**



3-(chloromethyl)benzoic acid **51**, gave thiomethyl compound **52** in 82% yield<sup>7</sup>. In a manner similar to that described for the preparation of compound **13** above, compound **52** was sequentially treated with gaseous chlorine to obtain the crude sulfonic acid **53** in theoretical yield followed by reaction with thionyl chloride which provided the crude acid chloride **54** as a waxy white solid. This was reacted directly  
 15 with anthranilate **21** to provide sufficiently pure sulfonyl chloride **55**, which was reacted with pyrrolidine to give a 26% yield of ester **56**. Subsequent hydrolysis with trifluoroacetic acid afforded the acid **57** in 83% yield as a white solid. **57**:  $^1\text{H}$  NMR

(DMSO-*d*<sub>6</sub>)  $\delta$  12.48 (s, 1 H), 8.86 (d,  $J = 7$  Hz, 1 H), 8.42 (d,  $J = 2$  Hz, 1 H), 8.12 (dd,  $J = 2, 7$  Hz, 1 H), 8.05 (s, 1 H), 7.95 (d,  $J = 6$  Hz, 1 H), 7.72 (d,  $J = 6$  Hz, 1 H), 7.64 (t,  $J = 6$  Hz, 1 H), 4.58 (s, 2 H), 3.20 (t,  $J = 5$  Hz, 4 H), 1.82 (m, 4 H) ppm.

- 5 **2-[(1,3-Benzoxazol-2-ylcarbonyl)amino]-5-cyanobenzoic acid** (36310-jcr-135a, PHA-734774, SPS# 0281864)



- To a solution of benzyl 1,3-benzoxazole-2-carboxylate (233 mg, 0.920 mmol) in 1:1  
 10 ethanol/THF (20 mL) was added palladium on carbon (56 mg of 5%, Aldrich) and triethylamine (180  $\mu$ L, 1.29 mmol, Aldrich). The mixture was stirred under 1 ATM of hydrogen for 2 hours and then filtered through a plug of celite. Removal of the solvent left the triethylamine salt as an orange oil (the protonated form of the acid rapidly decarboxylates and should be avoided). This oil was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (20  
 15 mL) and treated with DMF (20  $\mu$ L) followed by oxalyl chloride (220  $\mu$ L, 2.52 mmol, Aldrich). Solvent and excess oxalyl chloride were removed by rotary evaporation after 76 hours. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (20 mL), and benzyl 2-amino-5-cyanobenzoate (250 mg, 0.991 mmol) in pyridine (8 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of CH<sub>2</sub>Cl<sub>2</sub>.  
 20 This solution was washed with 2 X 100 of 1.0 M HCl and 100 mL of brine. Product was adsorbed onto silica gel and purified on a Biotage Flash 40 M silica gel cartridge using CH<sub>2</sub>Cl<sub>2</sub> as eluent. Product was collected as 218 mg of white solid as the benzyl ester. A mixture of benzyl 2-[(1,3-benzoxazol-2-ylcarbonyl)amino]-5-cyanobenzoate (168 mg, 0.423 mmol) and palladium on carbon (33 mg of 5%, Aldrich) in 2:1  
 25 THF/ethanol (30 mL) was stirred under 1 ATM of hydrogen for 25 minutes. The mixture was filtered through a plug of celite and then evaporated. The residue was dried at 100 °C under vacuum yielding 116 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*D*<sub>6</sub>)  $\delta$  ppm 7.56 (t,  $J = 7.67$  Hz, 1 H) 7.63 (t,  $J = 7.88$  Hz, 1 H) 7.94 (d,  $J = 8.29$  Hz, 1 H) 8.00 (d,  $J = 7.67$  Hz, 1 H) 8.16 (dd,  $J = 8.81, 1.97$  Hz, 1 H) 8.45 (d,  $J = 2.07$  Hz, 1 H) 8.87 (d,  $J = 8.71$  Hz, 1 H) 13.16 (s, 1 H).  
 30

The following compounds were produced via the methods described above using appropriate starting materials and making non-critical variations.

- 5 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)[1,1'-biphenyl]-3-carboxylic acid
- 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(2-furyl)benzoic acid
- 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(2-thienyl)benzoic acid
- 10 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(2-pyrazinyl)benzoic acid,
- 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(1-methyl-1H-pyrrol-2-yl)benzoic acid
- 15 4'-Chloro-4-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)[1,1'-biphenyl]-3-carboxylic acid
- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-3'-nitro[1,1'-biphenyl]-3-carboxylic acid
- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-4'-cyano[1,1'-biphenyl]-3-carboxylic acid
- 20 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(5-chloro-2-thienyl)benzoic acid
- 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(4-methyl-2-thienyl)benzoic acid
- 25 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-4'-fluoro[1,1'-biphenyl]-3-carboxylic acid
- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-2'-(trifluoromethyl)[1,1'-biphenyl]-3-carboxylic acid
- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-3',5'-bis(trifluoromethyl)[1,1'-biphenyl]-3-carboxylic acid
- 30 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(5-methyl-2-thienyl)benzoic acid

- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-2',4'-difluoro[1,1'-biphenyl]-3-carboxylic acid
- 4'-*t*-Butyl-4-({3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)[1,1'-biphenyl]-3-carboxylic acid
- 5 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-3'-(trifluoromethyl)[1,1'-biphenyl]-3-carboxylic acid
- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-4'-(trifluoromethyl)[1,1'-biphenyl]-3-carboxylic acid
- 4-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-2'-methyl[1,1'-biphenyl]-3-carboxylic acid
- 10

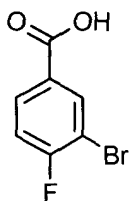
- 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(3,5-dimethyl-4-isoxazolyl)benzoic acid
- 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-(2,4-dimethoxy-5-pyrimidinyl)benzoic acid
- 5 2-[(3-[(4-Chlorophenyl)(methyl)amino]sulfonyl}benzoyl)amino]-5-(trifluoromethyl)benzoic acid,
- 2-[(3-Bromo-5-[(4-chlorophenyl)(methyl)amino]sulfonyl}benzoyl)amino]-5-chlorobenzoic acid
- 5-Bromo-2-[(3-[(4-chlorophenyl)(methyl)amino]sulfonyl}-5-nitrobenzoyl)amino]benzoic acid
- 10 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-cyanobenzoic acid
- 5-Bromo-2-({3-cyano-5-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl}amino}benzoic acid
- 15 5-Cyano-2-({3-(2,3-dihydro-1H-indol-1-ylsulfonyl)-5-methylbenzoyl}amino}benzoic acid
- Methyl 2-({3-[2-(acetyloxy)ethyl]-5-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl}amino}-5-cyanobenzoate
- 5-Cyano-2-({3-(2,3-dihydro-1H-indol-1-ylsulfonyl)-5-(2-hydroxyethyl)benzoyl}amino}benzoic acid
- 20 2-({3-Bromo-5-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl}amino}-5-chlorobenzoic acid
- 5-Chloro-2-[(3-[(4-chlorophenyl)(methyl)amino]sulfonyl}benzoyl)amino]benzoic acid
- 25 2-[(3-Bromo-5-[(4-chlorophenyl)(methyl)amino]sulfonyl}benzoyl)amino]-5-cyanobenzoic acid
- 5-cyano-2-[(3-[(2-hydroxyphenyl)(methyl)amino]sulfonyl}benzoyl)amino]benzoic acid
- 5-Bromo-2-[(5-[(4-chlorophenyl)(methyl)amino]sulfonyl}-2-methoxybenzoyl)amino]benzoic acid
- 30 5-Bromo-2-[(5-[(4-chlorophenyl)(methyl)amino]sulfonyl}-2-methylbenzoyl)amino]benzoic acid



- 5-Bromo-2-[(2-bromo-5-{{(4-chlorophenyl)(methyl)amino}sulfonyl} benzoyl)amino]benzoic acid
- 5-Bromo-2-[(3-{{(4-chlorophenyl)(methyl)amino}sulfonyl}-4-methoxybenzoyl)amino]benzoic acid
- 5 5-Bromo-2-[(3-{{(4-chlorophenyl)(methyl)amino}sulfonyl}-4-methylbenzoyl)amino]benzoic acid
- 5-Bromo-2-[(4-bromo-3-{{(4-chlorophenyl)(methyl)amino}sulfonyl} benzoyl)amino]benzoic acid
- 2-[(3-{{(4-Chlorophenyl)(methyl)amino}sulfonyl} benzoyl)amino]-5-nitrobenzoic acid
- 10 2-[(4-{{(4-Chlorophenyl)(methyl)amino}sulfonyl} benzoyl)amino]-5-nitrobenzoic acid
- 5-Bromo-2-[(3-{{(4-chlorophenyl)(methyl)amino}sulfonyl}-4-morpholin-4-ylbenzoyl)amino]benzoic acid
- 5-Bromo-2-[(3-bromo-5-{{(4-chlorophenyl)(methyl)amino}sulfonyl} benzoyl)amino] benzoic acid
- 15 2-{{[3-Bromo-5-(2,3-dihydro-1H-indol-1-ylsulfonyl)benzoyl]amino}-5-cyanobenzoic acid
- 2-{{[3-Bromo-5-(morpholin-4-ylsulfonyl)benzoyl]amino}-5-chlorobenzoic acid
- 5-Chloro-2-{{[3-(2,3-dihydro-1H-indol-1-ylsulfonyl)-5-methylbenzoyl]amino} benzoic acid
- 20 5-Iodo-2-{{[3-(morpholin-4-ylsulfonyl)benzoyl]amino} benzoic acid
- 2-{{[4-{{(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl} amino)-5-cyanobenzoic acid
- 2-{{[3-(Morpholin-4-ylsulfonyl)benzoyl]amino}-5-thiocyanatobenzoic acid

## 25 **Example 2: Amine, Ether, and Thioether Derivatives**

### **Preparation of 3-Bromo-4-fluorobenzoic acid**

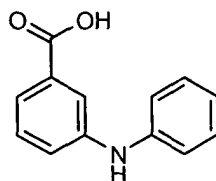


- 3-Bromo-4-fluoro-benzaldehyde (10.0 g, 49 mmol) in H<sub>2</sub>O(150 mL, followed by the
- 30 addition of KMnO<sub>4</sub> (15.5 g, 98 mmol) heated at reflux (foams extensively) for 1 h,

then added additional  $\text{KMnO}_4$  (15.5 g, 98 mmol) and continued heating for another 3 h. The reaction was cooled to rt, then filtered through Celite. The solution was acidified with HCl, and the resulting white precipitate was filtered off, to afford 6.1 g (56%) of a white solid.

5

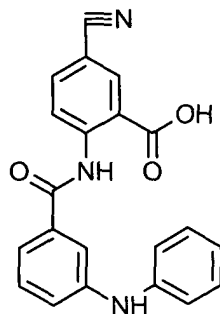
### Preparation of 3-Anilinobenzoic acid



Methyl 3-bromobenzoate (1000 mg, 4.65 mmol),  $\text{Pd}_2(\text{dba})_3$  (53 mg, 0.058 mmol),  $\text{Cs}_2\text{CO}_3$  (2120 mg, 1.4 mmol) and N-[2'-(dicyclohexylphosphino)-1,1'-biphenyl-2-yl]-N,N-dimethylamine (27mg, 0.07 mmol) were placed in a 100ml one-necked round bottom flask. The system was evacuated and filled with argon several times. Then aniline (519 mg, 5.58 mmol) was added, followed by the addition of toluene (50 ml). The solution was heated at  $100^\circ\text{C}$  for 20h, the solvent was removed in vacuo and residue was purified by silica gel chromatography (EtOAc/heptane 1/3) to get 180 mg (18%) of methyl ester as a yellow solid, which was hydrolyzed by LiOH (50 mg) in THF (4 ml) and water (1 ml) to afford 140 mg (82%) of **3-Anilinobenzoic acid** as a white solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  8.02 (s, 1 H), 7.65 (s, 1 H), 7.33 (d,  $J$  = 7.5 Hz, 1 H), 7.19 (t,  $J$  = 8.3 Hz, 2 H), 7.10 (d,  $J$  = 7.7 Hz, 1H), 7.03 (d,  $J$  = 7.6 Hz, 2 H), 6.96 (m, 1 H), 6.76 (t,  $J$  = 7.3 Hz, 1 H);

20

### 2-[(3-Anilinobenzoyl)amino]-5-cyanobenzoic acid



Prepared according to the general methods described above: **3-Anilinobenzoic acid** (140 mg, 0.66 mmol) and PHA-561053 (130 mg, 0.59 mmol) afforded 61 mg (25%)

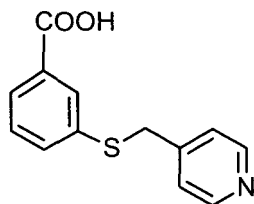
25

of t-butyl ester as a yellow solid, which was hydrolyzed to 48 mg (91%) of a green solid.

**Analytical data for PHA-610938**

<sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.81 (d, *J* = 9.0 Hz, 1 H), 8.46 (s, 1 H), 8.35 (d, *J* = 2.2 Hz, 1 H), 7.82 (dd, *J* = 1.9, 8.8 Hz, 1 H), 7.72 (s, 1 H), 7.42 (m, 2 H), 7.27 (m, 3 H), 7.14 (d, *J* = 7.8 Hz, 2 H), 6.88 (t, *J* = 7.3 Hz, 1 H).

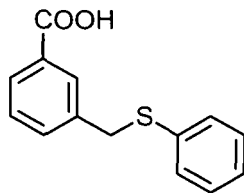
**Preparation of 3-[(Pyridin-4-ylmethyl)thio]benzoic acid**



10

Water (10 mL) was added to a flask containing 3-mercaptopbenzoic acid (2.08 g, 13.5 mmol, Aldrich) and sodium hydroxide (1.16 g, 29.0 mmol). To the resulting solution was added 4-picolyl chloride hydrochloride (2.31 g, 14.1 mmol, Aldrich) and ethanol (20 mL). The mixture was heated in a 75 °C oil bath for 1 hour and then added to a separatory funnel with 100 mL of water and 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. This resulted in a suspension in the aqueous layer. This suspension was washed with an additional 100 mL of CH<sub>2</sub>Cl<sub>2</sub> and then filtered. The solid was then dried at 100 °C under vacuum yielding 2.80 g of white solid.

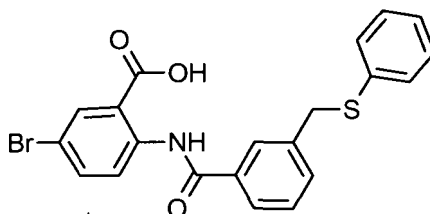
**Preparation of 3-[(Phenylthio)methyl]benzoic acid**



20

To a solution of the corresponding methyl ester described by Holoboski, M.A.; Koft, E. in *J. Org. Chem.*, **1992**, 57, 965-969, (1.23 g, 4.76 mmol) in methanol (15 mL) was added 1.0 M aqueous NaOH (8.0 mL). The resulting mixture was heated in a 50 °C oil bath for 1.5 hours. Most of the methanol was removed by rotary evaporation, and the residue was added to a separatory funnel with 100 mL of 1.0 M aqueous HCl and 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The CH<sub>2</sub>Cl<sub>2</sub> was washed with another 100 mL of 1.0 M aqueous HCl followed by 100 mL of water and then dried over Na<sub>2</sub>SO<sub>4</sub>. Solvent was removed, and the residue was dried at 100 °C yielding 1.11 g of white solid.

**5-Bromo-2-({3-[(phenylthio)methyl]benzoyl}amino)benzoic acid**



To 3-[(phenylthio)methyl]benzoic acid (400 mg, 1.64 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (15 mL) was added DMF (20 µL) and oxalyl chloride (200 µL, 2.29 mmol). The mixture was stirred for 1.5 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (15 mL), and methyl 2-amino-5-bromobenzoate (330 mg, 1.43 mmol, Avocado) in pyridine (8 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The CH<sub>2</sub>Cl<sub>2</sub> was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 50% CH<sub>2</sub>Cl<sub>2</sub>/heptane to 75% CH<sub>2</sub>Cl<sub>2</sub>/heptane as eluent. Yield was 544 mg of white solid as the methyl ester.

To a mixture of the corresponding methyl ester (386 mg, 0.845 mmol) in dioxane (20 mL) was added 1 M aqueous sodium hydroxide (2.0 mL). The mixture was stirred for at room temperature for 1.25 hours and then at 50 °C for 1.5 hours. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The CH<sub>2</sub>Cl<sub>2</sub> was washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of brine. It was then dried over Na<sub>2</sub>SO<sub>4</sub> and evaporated. The residue was recrystallized from hot ethanol (8

mL). The solids were washed with ethanol followed by heptane and then dried at 100 °C under vacuum yielding 279 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.08 (s, 1 H), 8.64 (d, *J* = 9.2 Hz, 1 H), 8.12 (d, *J* = 2.5 Hz, 1 H), 7.97 (s, 1 H), 7.86 (dd, *J* = 9.2, 2.5 Hz, 1 H), 7.80 (d, *J* = 7.6 Hz, 1 H), 7.61 (d, *J* = 7.6 Hz, 1 H), 7.51 (t, *J* = 7.6 Hz, 1 H), 7.35 (d, *J* = 7.1 Hz, 2 H), 7.29 (t, *J* = 7.9 Hz, 2 H), 7.18 (t, *J* = 7.1 Hz, 1 H), 7.35 (s, 2 H).

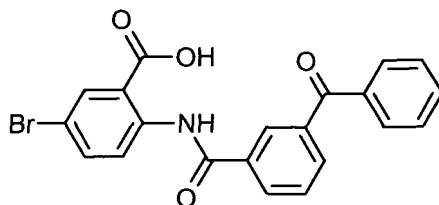
Other compounds produced via the above-described methodology using appropriate starting materials and making non-critical variations include:

- 10 2-{{3-(benzylthio)benzoyl}amino}-5-bromobenzoate
- 2-{{3-(Benzyloxy)benzoyl}amino}-5-bromobenzoic acid
- 5-Bromo-2-{{3-(ethylthio)benzoyl}amino}benzoic acid
- Methyl-5-Bromo-2-({3-[(pyridin-4-ylmethyl)thio]benzoyl}amino)benzoate
- 5-Bromo-2-({3-[(pyridin-4-ylmethyl)thio]benzoyl}amino)benzoic acid
- 15 5-bromo-2-({3-[(pyridin-4-ylmethyl)thio]benzoyl}amino)benzoic acid hydrochloride
- 5-Bromo-2-[(3-phenoxybenzoyl)amino]benzoic acid
- 5-Bromo-2-{{3-(phenylthio)benzoyl}amino}benzoic acid
- 5-Cyano-2-[(3-phenoxybenzoyl)amino]benzoic acid
- 5-Cyano-2-({3-[(pyridin-4-ylmethyl)thio]benzoyl}amino)benzoic acid
- 20 5-Cyano-2-({3-[(pyridin-4-ylmethyl)thio]benzoyl}amino)benzoic acid hydrochloride
- 2-{{3-(Benzyloxy)benzoyl}amino}-5-cyanobenzoic acid
- 2-{{3-(Benzylthio)benzoyl}amino}-5-cyanobenzoic acid
- 5-cyano-2-({3-[(1-phenylethyl)thio]benzoyl}amino)benzoic acid
- 5-cyano-2-{{3-(cyclopentylthio)benzoyl}amino}benzoic acid
- 25 5-cyano-2-{{3-(cyclopentylsulfinyl)benzoyl}amino}benzoic acid
- 5-Chloro-2-[(4-methoxy-3-nitrobenzoyl)amino]benzoic acid
- 2-{{4-(Benzylsulfanyl)-3-bromobenzoyl}amino}-5-chlorobenzoic acid
- 5-Cyano-2-{{3-(3-fluorophenoxy)benzoyl}amino}benzoic acid
- 5-Cyano-2-{{3-(2-methylphenoxy)benzoyl}amino}benzoic acid
- 30 5-Cyano-2-{{3-(4-methoxyphenoxy)benzoyl}amino}benzoic acid
- 5-Cyano-2-{{3-(3-nitrophenoxy)benzoyl}amino}benzoic acid

### Example 3: KETONE DERIVATIVES

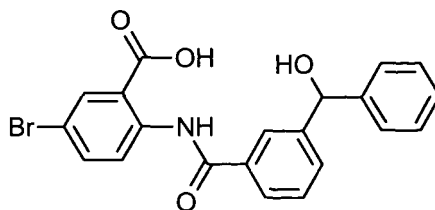
#### 2-[(3-Benzoylbenzoyl)amino]-5-bromobenzoic acid

5



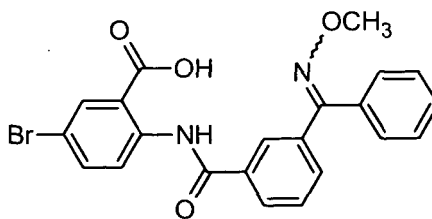
To 3-benzoylbenzoic acid (633 mg, 2.80 mmol, Aldrich) in  $\text{CH}_2\text{Cl}_2$  (20 mL) was added DMF (20  $\mu\text{L}$ ) and oxalyl chloride (450  $\mu\text{L}$ , 5.16 mmol). The mixture was stirred for 1.7 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (20 mL), and methyl 2-amino-5-bromobenzoate (565 mg, 2.46 mmol, Avocado) in pyridine (6 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of  $\text{CH}_2\text{Cl}_2$ . This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 75%  $\text{CH}_2\text{Cl}_2$ /heptane to 100%  $\text{CH}_2\text{Cl}_2$  as eluent. Yield was 825 mg of white solid as the methyl ester. To a mixture of the corresponding methyl ester (645 mg, 1.47 mmol) in dioxane (20 mL) was added 1 M aqueous sodium hydroxide (3.0 mL). The mixture was stirred in a 50  $^\circ\text{C}$  oil bath for 2 hours. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of  $\text{CH}_2\text{Cl}_2$ . The organics were washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. They were then dried over  $\text{MgSO}_4$  and evaporated. The residue was recrystallized from hot ethanol/THF. The solids were washed with ethanol followed by pentane and then dried at 100  $^\circ\text{C}$  under vacuum yielding 329 mg of white solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.17 (s, 1 H), 8.61 (d,  $J$  = 9.2 Hz, 1 H), 8.31 (s, 1 H), 8.23 (d,  $J$  = 7.6 Hz, 1 H), 8.12 (d,  $J$  = 2.0 Hz, 1 H), 7.99 (d,  $J$  = 7.6 Hz, 1 H), 7.87 (dd,  $J$  = 9.2, 2.5 Hz, 1 H), 7.77-7.82 (m, 3 H), 7.73 (t,  $J$  = 7.4 Hz, 1 H), 7.61 (t,  $J$  = 7.6 Hz, 2 H).

**5-Bromo-2-({3-[hydroxy(phenyl)methyl]benzoyl}amino)benzoic acid**



Solid sodium borohydride (82 mg, 2.2 mmol) was added in one portion to a slurry of methyl 2-[(3-benzoylbenzoyl)amino]-5-bromobenzoate (826 mg, 1.88 mmol) in 40 mL of 1:1 methanol/THF. The mixture was stirred for 75 minutes before being quenched by the addition of 1 M aqueous HCl (50 mL). The organics were removed by rotary evaporation, and the product was extracted into 100 mL + 50 mL of CH<sub>2</sub>Cl<sub>2</sub>. The CH<sub>2</sub>Cl<sub>2</sub> was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from CH<sub>2</sub>Cl<sub>2</sub> to 5% EtOAc/CH<sub>2</sub>Cl<sub>2</sub> as eluent. Yield was 433 mg of white solid as the methyl ester. To a mixture of the corresponding methyl ester (348 mg, 0.788 mmol) in dioxane (20 mL) was added 1 M aqueous sodium hydroxide (1.5 mL). The mixture was stirred at room temperature overnight and then heated in a 50 °C oil bath for 30 minutes. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The organics were washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. They were then dried over MgSO<sub>4</sub> and evaporated. The residue was recrystallized from hot ethanol (10 mL). The solids were washed with ethanol followed by pentane and then dried at 100 °C under vacuum yielding 130 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.12 (s, 1 H), 8.66 (d, *J* = 8.7 Hz, 1 H), 8.13 (d, *J* = 2.5 Hz, 1 H), 8.05 (s, 1 H), 7.85 (dd, *J* = 9.2, 2.5 Hz, 1 H), 7.79 (d, *J* = 7.6 Hz, 1 H), 7.62 (d, *J* = 8.1 Hz, 1 H), 7.51 (t, *J* = 7.6 Hz, 1 H), 7.42 (d, *J* = 7.1 Hz, 2 H), 7.32 (t, *J* = 7.6 Hz, 2 H), 7.22 (t, *J* = 7.1 Hz, 1 H), 6.07 (br s, 1 H), 5.81 (s, 1 H).

**5-Bromo-2-({3-[(methoxyimino)(phenyl)methyl]benzoyl}amino)benzoic acid (PHA-522146)**

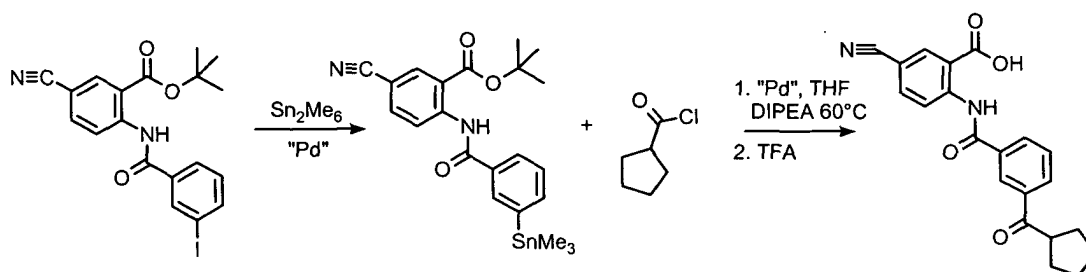


00055 US1  
Methyl 2-[(3-benzoylbenzoyl)amino]-5-bromobenzoate (763 mg, 1.74 mmol) was dissolved in 60 mL of 1:1 EtOH/pyridine with warming. After this solution was allowed to cool, solid O-methylhydroxylamine hydrochloride (350 mg, 4.19 mmol, Aldrich) was added in one portion. The resulting slurry was stirred at room

5 temperature for 6 days, after which it was a solution. The solvents were removed by rotary evaporation, and the residue was dissolved in 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The CH<sub>2</sub>Cl<sub>2</sub> was dried over MgSO<sub>4</sub> and evaporated leaving 785 mg of white solid that was approximately a 1:1 mixture of oxime isomers by <sup>1</sup>H NMR. To a mixture of the  
10 corresponding methyl ester (470 mg, 1.01 mmol) in dioxane (15 mL) was added 1 M aqueous sodium hydroxide (2.0 mL). The mixture was stirred at room temperature overnight. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The organics were washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of  
15 water. They were then dried over MgSO<sub>4</sub> and evaporated. The orange residue was recrystallized from hot ethanol (10 mL). The solids were washed with ethanol followed by heptane and then dried at 100 °C under vacuum yielding 255 mg of white solid that was approximately a 1:1 mixture of oxime isomers by <sup>1</sup>H NMR. Due to the presence of 2 isomers, the NMR is difficult to assign. At 400 MHz in DMSO-*d*<sub>6</sub>, the  
20 amide protons appear as singlets at 12.10 and 12.07 ppm. The aromatic protons appear between 7.32 and 8.63 ppm. The methyl peaks appear as singlets at 3.93 and 3.92 ppm.

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**5-cyano-2-[[3-(cyclopentylcarbonyl)benzoyl]amino]benzoic acid**





tert-Butyl 5-cyano-2-[(3-iodobenzoyl)amino]benzoate (1.0 g, 2.23 mmol) was dissolved in 20 ml of CH<sub>2</sub>Cl<sub>2</sub>. Hexamethylditin (1.1 g, 3.35 mmol) and allylpalladium chloride dimer (73 mg, 0.2 mmol) were then added and the mixture stirred at room temperature for 5 hr. The reaction was diluted with CH<sub>2</sub>Cl<sub>2</sub> then washed with water.

- 5 The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The remaining oil was purified via silica gel chromatography to give 670 mg (62%) of the desired tin compound. This product was subsequently dissolved in 15 mL of THF. To this was added DIPEA (1 mL), Pd<sub>2</sub>dba<sub>3</sub> (115 mg, .125 mmol) and cyclopentanecarbonyl chloride (230 mg 1.73 mmol). The reaction was then warmed
- 10 to 60 °C and stirred for 10 additional hr. After cooling to room temperature the reaction was poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The remaining residue was purified via silica gel chromatography, giving 415 mg (72%) of the desired ketone. The ketone was treated with CH<sub>2</sub>Cl<sub>2</sub>/TFA and stirred for 10
- 15 additional hours. The solvent was removed *in vacuo* and the remaining solid was recrystallized from MeOH to give the title compound (329 mg, 91%) as a white solid. <sup>1</sup>H NMR (400 MHz, DMSO) 1.62-1.67 (m, 4H), 1.73-1.80 (m, 2H), 1.92-1.98 (m, 2H), 3.90 (quint, 1H), 7.77 (t, 1H), 8.11 (dd, 1H), 8.19 (d, 1H), 8.27 (d, 1H), 8.41 (d, 1H), 8.53 (s, 1H), 8.84 (d, 1H), 12.55 (s, 1H)

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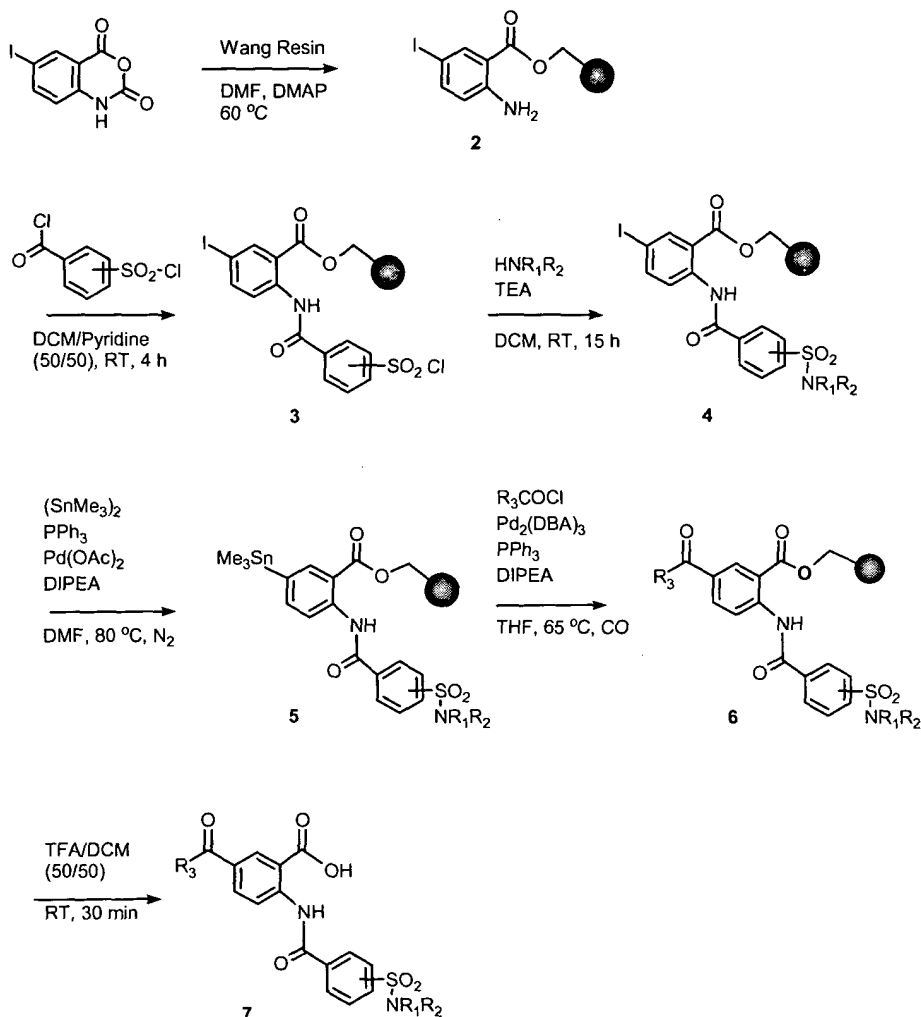
Other compounds produced via the above-described methodology using appropriate starting materials and making non-critical variations include:

- 2-[(3-Benzoylbenzoyl)amino]-5-chlorobenzoic acid
- 2-[(4-Acetylbenzoyl)amino]-5-bromobenzoic acid
- 25 2-[(4-Benzoylbenzoyl)amino]-5-bromobenzoic acid
- 2-[(3-Acetylbenzoyl)amino]-5-bromobenzoic acid
- 5-Bromo-2-({3-[(hydroxyimino)(phenyl)methyl]benzoyl}amino)benzoic acid
- (+)-5-Bromo-2-({3-[hydroxy(phenyl)methyl]benzoyl}amino)benzoic acid
- (-)-5-bromo-2-({3-[hydroxy(phenyl)methyl]benzoyl}amino)benzoic acid
- 30 **2-[(3-Benzoylbenzoyl)amino]-5-nitrobenzoic acid**
- 2-[(3-Benzoylbenzoyl)amino]-5-cyanobenzoic acid**
- 5-Cyano-2-({3-[(hydroxyimino)(phenyl)methyl]benzoyl}amino)benzoic acid**
- 5-Cyano-2-({3-[(methoxyimino)(phenyl)methyl]benzoyl}amino)benzoic acid**

## Solid Phase Synthesis

Additional methodologies for producing compounds of this invention are shown below.

### 5 Scheme 3.1

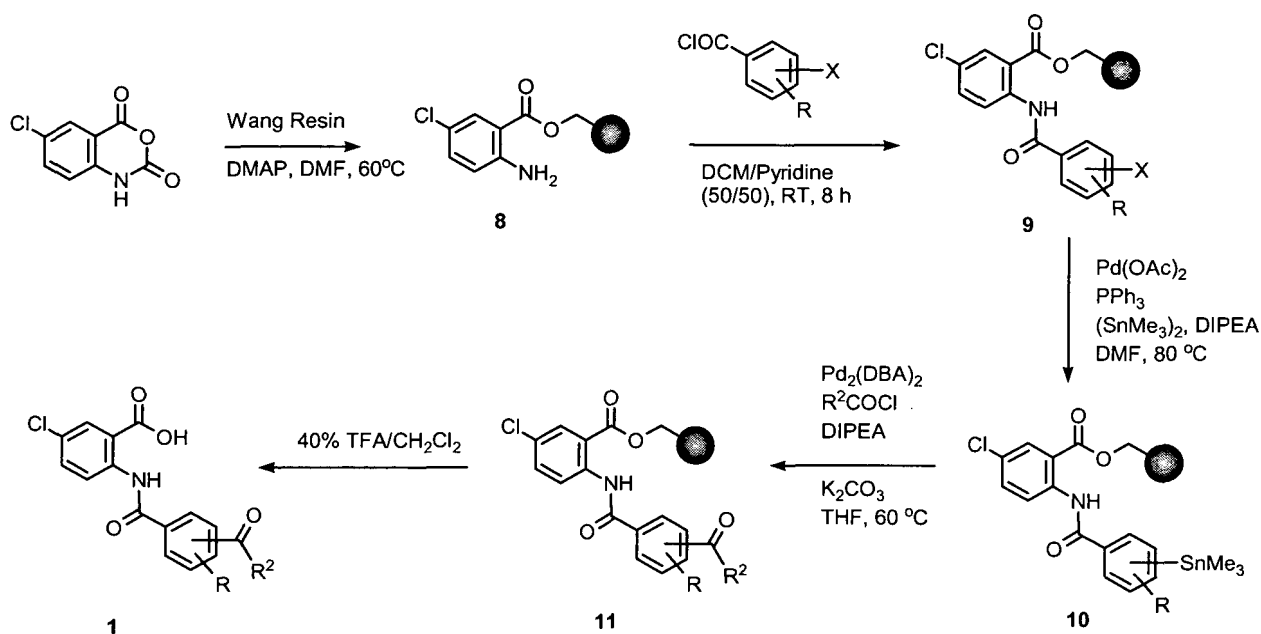


$\text{R}_3$  is a  $\text{C}_{1-4}$ alkyl optionally substituted with 1-3 halo, -OH,  $\text{NO}_2$ , or -CN.

Development of a solid phase route to ketones **1** was effected by a similar route and is summarized in Scheme 3.2. Chlorine was selected as the anthranilic acid 5-substituent instead of the 5-bromine of the ketone leads in order to avoid the potential for competing reactions in the ensuing palladium-catalyzed stannylation. Solid-supported aryl halide **8** was prepared by reaction of chloroisatoic anhydride with Wang resin. Coupling with halo ( $\text{X} = \text{Br}$  or  $\text{I}$ ) aryl chlorides then afforded

benzamides **9**, which were stannylated with hexamethyl distannane under the influence of palladium catalyst using the same conditions that were applied in Scheme 3.1. The subsequent carbonylation reactions were found to be optimal using the slightly modified conditions of Ellman.<sup>8</sup> Eliminating the ligand altogether and adding potassium carbonate as another proton scavenger slightly enhanced the rate of the reactions and the product purities in the end. Carbon monoxide was not necessary to eliminate aryl-aryl coupling by-products. One other modification in the synthetic conditions was to decrease the amount of TFA used in the cleavage cocktail in order to avoid trace amounts of a cleavage impurity.

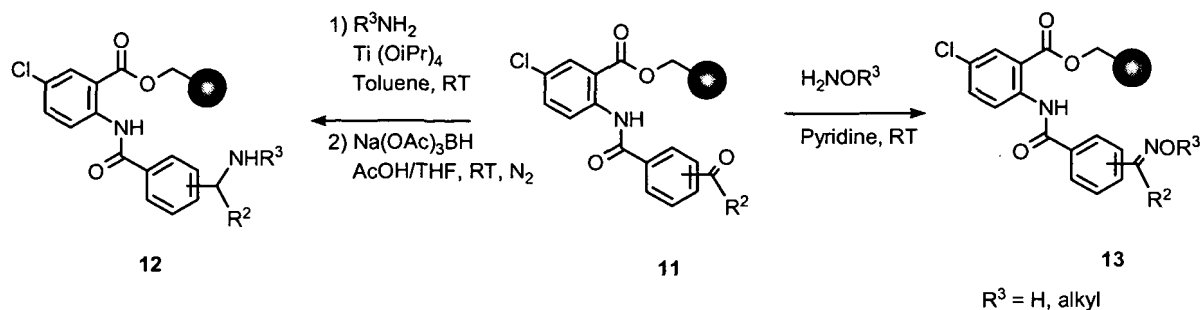
**Scheme 3.2**



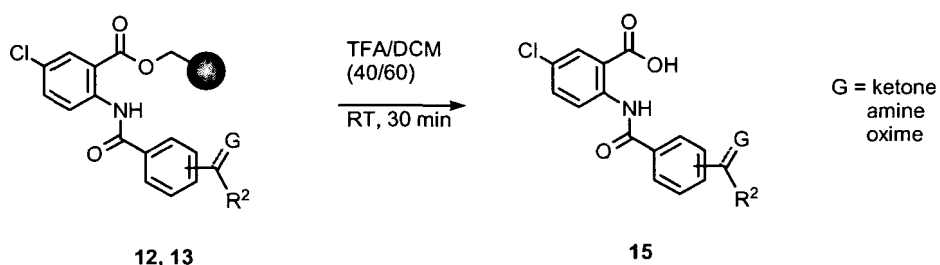
### Generation of Oximes and Amines from Solid-Supported Ketones

Chemistry was developed for amine (**12**) and oxime (**13**) derivatization of the ketones on solid-phase (Scheme 3.3). Following TFA cleavage (Scheme 3.4), the amines could be successfully purified by trapping the products on sulfonic acid resin and then washing off with 2 N NH<sub>4</sub>OH/methanol. The remaining compounds were subjected to preparative HPLC purification.

**Scheme 3.3**



**Scheme 3.4**



5

## Ketones

### Step 1: Preparation of 8

- 10 To 4.5 grams of Wang resin (Irori Unisphere, 1.36 mmol/g loading, 6.12 mmol) in a 125 mL serum bottle, 60 mL of DMF were added followed by 6.1 grams (5 eq., 30.6 mmol) of 5-chloroisotoic anhydride and 3.74 grams (5 eq., 30.6 mmol) of 4-dimethylaminopyridine. The serum bottle was purged with nitrogen, capped, and shaken on an orbital mixer at 60 °C. Initially, the reagent cocktail was not
- 15 homogeneous, but after several hours, a concentrated solution had formed around the swelled resin. After 18 hours, the reaction slurry was cooled and transferred to a 60 mL syringe-barrel reaction vessel. The reagent cocktail was then drained and the resin washed as follows: 3 X (acetonitrile, DMF), then 3 X (acetonitrile, methylene chloride). The resin was treated a second time with 60 mL of DMF, 6.1 grams (5 eq.,
- 20 30.6 mmol) of 5-chloroisotoic anhydride, and 3.74 grams (5 eq., 30.6 mmol) of 4-dimethylaminopyridine. Following mixing at 60 °C for 6 hours, the reagent cocktail was again drained and the resin washed as above. In a vacuum oven at 25 °C, the resin was dried for 72 hours to give a final weight of 5.36 grams (1.14 mmol/g loading).

## Step 2: Preparation of 9

To 6.7 mmol of the halo benzoic acids suspended in 20 mL of methylene chloride, 20  $\mu$ L of DMF and 1.17 mL (1.7 grams, 13.4 mmol, 2 eq.) of oxalyl chloride were added.

5 The flasks were sealed and stirred with occasional release of gas build-up. After stirring overnight, the reaction mixtures had become almost completely homogeneous with no more gas build-up. Solvent and excess oxalyl chloride were then evaporated *in vacuo* to dryness. The acid chlorides were re-dissolved in 10 mL of methylene chloride and added to 1 gram of resin 8 (1.14 mmol/gram loading, 1.14 mmol)

10 swollen with 10 mL of pyridine in 25 mL vials. Some fuming was observed initially. The mixtures were purged with nitrogen for 10 seconds then the vials capped, and the mixtures shaken at room temperature for 4 hours. By that time, the resins had taken on a light orange color and a tan precipitate had formed in the supernatant. The reagent solutions were then drained in syringe-barrel reaction vessels and the resins

15 rinsed five times with alternating acetonitrile and methylene chloride washes. The resins were kept wet with methylene chloride until used in the next step. Cleavage aliquots (40 % TFA/CH<sub>2</sub>Cl<sub>2</sub>) had purities of > 80% by HPLC and were registered as PHA compounds (Table 1).

## 20 Step 3: Preparation of 10

A stock solution of palladium acetate (0.1 eq., 0.01 mmol, 0.0022 g per 1 mL), triphenylphosphine (0.25 eq., 0.025 mmol, 0.0065 g per 1 mL), and diisopropylethylamine (0.5 eq., 0.05 mmol, 0.0065 g, 0.0087 mL per 1 mL) in 6.5 mL DMF (degassed with N<sub>2</sub>) was prepared. To each of the resins (9) in 8 mL vials, 1 mL

25 of stock catalyst solution was added, followed by 0.042 mL of hexamethyl ditin (2.0 eq., 0.2 mmol, 0.065 g). Each vial was purged with nitrogen and then capped. The reaction mixtures were then heated to 60 °C and mixed in an orbital shaker for 17 h. By that time, the resins had all turned black in color. Following cooling, the reaction mixtures were transferred to filter vessels, and reagents were drained. This was

30 followed by washing three times with DMF, three times with alternating acetonitrile/DMF, three times with alternating acetonitrile/methylene chloride, and twice with THF. Cleavage aliquots were taken (cleaved in 40/60 TFA/CH<sub>2</sub>Cl<sub>2</sub>) to check for completion of reaction by monitoring the protodestannylation products.

#### Step 4: Preparation of 11

To each of the 8 mL vials holding resins **10**, 2 mL of a THF (degassed with carbon monoxide) stock solution containing: 0.0046 g of tris (dibenzylidene acetone)  
5 dipalladium (0) (0.05 eq., 0.005 mmol, per 2 mL THF); 0.0052 g of triphenylphosphine (0.2 eq, 0.02 mmol, per 2 mL THF); and 0.139 mL diisopropyl ethylamine (8 eq., 0.80 mmol, 0.103 g, 0.139 mL per 2 mL) were added. Commercially available acid chlorides (8 eq., 0.8 mmol) were then added. The reaction vessels were purged with carbon monoxide, capped and shaken at 60 °C for  
10 18 h. When cool, the reaction mixtures were filtered through fritted syringe barrels, then the resins rinsed six times with alternating acetonitrile/methylene chloride washes and dried under vacuum at room temperature.

#### Step 5: Preparation of 1

15 To each of the fritted vessels containing resins **11**, 2 mL of the cleavage cocktail (40/60 TFA/CH<sub>2</sub>Cl<sub>2</sub>) were added and the mixtures swirled for 45 minutes. Cleavage filtrates were then collected in tared vials followed by stripping of solvents *in vacuo*. The residues were analyzed by HPLC and ESMS separately. The library was then purified by preparative HPLC. Results for the library both pre- and post-purification  
20 are compiled in Table 5.

#### Preparation of Oximes 13

Ketone precursors to the oxime derivatives were produced as shown above. To 0.1 gram (~0.12 mmol) of the ketone resins **11** in a 48 well Robbins Block, 2 mL of  
25 pyridine were added followed by 10 equivalents (1.2 mmol) of each alkoxyamine (hydroxylamine hydrochloride; methoxyamine hydrochloride; o-benzyloxyamine hydrochloride; and o-allylhydroxylamine hydrochloride). The reaction block was sealed and mixed overnight at room temperature in the rotating oven. After 20 hours, the resins resins were drained and washed with 3 X (MeOH,  
30 CH<sub>2</sub>Cl<sub>2</sub>) and 3 X (MeCN, CH<sub>2</sub>Cl<sub>2</sub>). Methanol was used early in the wash cycle because MeCN and CH<sub>2</sub>Cl<sub>2</sub> left a precipitate in the supernatant at that point. Treatment of the resins with 40 % TFA/CH<sub>2</sub>Cl<sub>2</sub> for 45 minutes afforded crude

products. Four of the library compounds (shown in Table 3) were then successfully purified (>90 % pure) via LC/MS.

### **Amine Derivatives**

#### **Preparation of Amines 12**

Into four 8 mL vials containing 0.1 grams (~0.12 mmols) of ketone resin **11**, 1.5 mL of toluene along with 0.12 grams (0.42 mmol) of titanium isopropoxide and 2.5 equivalents (0.30 mmol) of each respective amine were added. The vials were purged with nitrogen, sealed with teflon-lined caps, and mixed at room temperature for 16 hours on an orbital shaker. At that time, 0.5 mL of THF, 0.1 mL of acetic acid, and 0.24 grams (1.14 mmol) of sodium triacetoxyborohydride were added and the slurry was mixed at room temperature. After 4 hours, the reagents were drained and the resin washed: 3 X (MeOH, DMF), 4X (MeOH, CH<sub>2</sub>Cl<sub>2</sub>). Treatment of the resin with 40 % TFA/CH<sub>2</sub>Cl<sub>2</sub> for 45 minutes afforded crude products in the purities included in Table 6. Crude product identities were confirmed by ES/MS.

#### **Step 1: Preparation of 8**

To 10.0 grams of Wang resin (Irori Unisphere, 1.36 mmol/g loading, 13.6 mmol) in a 250 mL serum bottle, 90 mL of DMF were added followed by 13.4 grams (5 eq., 68 mmol) of 5-chloroisotoic anhydride and 8.3 grams (5 eq., 68 mmol) of 4-dimethylaminopyridine. The serum bottle was purged with nitrogen, capped, and shaken on an orbital mixer at 60 °C. Initially, the reagent cocktail was not homogeneous, but after several hours, a concentrated solution had formed around the swelled resin. After 18 hours, the reaction slurry was cooled and transferred to a 60 mL syringe-barrel reaction vessel. The reagent cocktail was then drained and the resin washed as follows: 3 X (acetonitrile, DMF), then 3 X (acetonitrile, methylene chloride). The resin was treated a second time with 90 mL of DMF, 13.4 grams (5 eq., 68 mmol) of 5-chloroisotoic anhydride, and 13.4 grams (5 eq., 68 mmol) of 4-dimethylaminopyridine. Following mixing at 60°C for 6 hours, the reagent cocktail was again drained and the resin washed as above. In a vacuum oven at 25°C, the resin was dried for 72 hours to give a final weight of 10.46 grams (1.30 mmol/g loading).

#### **Step 2: Preparation of 9 (R = H)**

To 6.2 grams (25 mmol) of the meta- and para- iodo benzoic acids suspended in 70 mL of methylene chloride, 40  $\mu$ L of DMF and 4.4 mL (6.35 grams, 50 mmol, 2 eq.) of oxalyl chloride were added. The serum bottles were sealed and stirred with occasional release of gas build-up. After stirring for 5 hours, the reaction mixtures had become almost completely homogeneous with no more gas build-up. Solvent and excess oxalyl chloride were then evaporated *in vacuo* to dryness. The acid chlorides were re-dissolved in 30 mL of methylene chloride and added to 4 gram of resin **8** (1.30 mmol/gram loading, 5.2 mmol) swollen with 30 mL of pyridine in 125 mL serum bottles. Some fuming was observed initially. The mixtures were purged with nitrogen for 10 seconds then the vials capped, and the mixtures shaken at room temperature for 4 hours. By that time, the resins had taken on a light orange color and a tan precipitate had formed in the supernatant. The reagent solutions were then drained in syringe-barrel reaction vessels and the resins rinsed five times with alternating acetonitrile and methylene chloride washes. The resins were then dried *in vacuo* to afford 5.14 g of the meta-iodo product and 5.09 g of the para-iodo product. Cleavage aliquots were >95 % pure by HPLC, with their identities confirmed by ESMS.

### Step 3: Preparation of **10** (R = H)

A stock solution of palladium acetate (0.012 M), triphenylphosphine (0.03 M), and diisopropylethylamine (0.06 M) in 80 mL DMF (degassed with N<sub>2</sub>) was prepared. To 4.0 grams (~5.0 mmol) of each resin (**9**) in 125 mL serum bottles, 40 mL of the stock catalyst solution were added, followed by 2.0 mL of hexamethyl ditin (2.0 eq., 9.6 mmol, 3.14 g). Each bottle was purged with nitrogen and then capped. The reaction mixtures were then heated to 60 °C and mixed in an orbital shaker for 17 h. By that time, the two resins had turned black in color. Following cooling, the reaction mixtures were transferred to filter vessels, and reagents were drained. This was followed by washing three times with DMF, three times with alternating acetonitrile/DMF, three times with alternating acetonitrile/methylene chloride, and twice with THF. Cleavage aliquots were taken (cleaved in 40/60 TFA/CH<sub>2</sub>Cl<sub>2</sub>) to check for completion of reaction by monitoring the protodestannylation products. Following cleavage, the meta-substituted resin gave 87 % of the expected destannylated product by HPLC, while the para-substituted isomer gave 70 %. Little



to no iodide starting material remained. The major impurity in both cases was an unidentified peak with  $[M+H]^+ = 369$  m/z.

#### Step 4: Preparation of 11

5 To each carboxylic acid weighed into a 20 mL vial (2.88 mmol), 6.5 mL of THF, 10  $\mu$ L of DMF, and 0.293 ml of oxalyl chloride (0.95 eq., 2.7 mmol, 3.35 g) were added. The vials were sealed and reaction mixtures shaken at room temperature for 4 hours with occasional release of evolved gas. In the meantime, the two stannylated resins (10) were distributed into Irori minikans (60 mg per kan), and the 72 kans were then  
10 distributed into twelve 125 mL serum bottles (six kans per bottle). To each of the bottles, 20 mL of a nitrogen degassed THF stock solution containing: tris (dibenzylidene acetone) dipalladium (0) (0.001 M); potassium carbonate (0.02 M); and diisopropyl ethylamine (0.10 M) were added. The THF solutions of acid chlorides (2.88 mmol, 6 eq.) were then added to their respective set of six bottles. The  
15 capped reaction vessels were purged with nitrogen, degassed, and shaken at 65 °C for 18 h. When cool, the resin containing kans were rinsed five times with alternating acetonitrile/ methylene chloride washes and dried under vacuum at room temperature. A cleavage aliquot revealed that ketone formation had gone to completion.

#### 20 Step 5a: Preparation of Resin-Bound Amines 12

To a 125 mL serum bottle containing 24 Irori cans loaded with resin 11, 30 mL of toluene were added, followed by 1.23 grams (6.0 mmol, 3.5 eq.) of titanium isopropoxide and 0.25 grams (4.3 mmol, 2.5 eq.) of propyl amine. The bottle was  
25 degassed to remove air bubbles from the Irori kans, then purged with nitrogen, sealed and mixed for 17 hours at room temperature. At that time, 10 mL of toluene, 2 mL of acetic acid, and 3.5 grams (16.3 mmol, 9.5 eq.) of sodium triacetoxy borohydride were added, and bottle re-purged and sealed, and mixed for 14 hours. Reagents were then drained and the resins washed three times with methanol and five times with  
alternating methanol/methylene chloride.

30

#### Step 5b: Preparation of Resin-Bound Oximes 13

To a 125 mL serum bottle containing 24 Irori kans loaded with resin 11, 40 mL of pyridine were added followed by 1.2 grams (17.2 mmols, 10 eq.) of hydroxylamine

hydro chloride. . The bottle was degassed to remove air bubbles from the Irori kans, then purged with nitrogen, sealed and mixed for 17 hours at room temperature. At that time, reagents were drained and the resins were washed three times with methanol, and five times with alternating methanol/methylene chloride.

5

#### Step 6: Preparation of 15

The 72 kans containing resins **12,13** were distributed into tared 8 mL vials and treated with 3 mL of TFA/CH<sub>2</sub>Cl<sub>2</sub> (40/60). The vials were degassed, capped, and mixed at room temperature for 1.5 hours. The kans were then plucked out of the vials using a  
10 syringe needle and washed with another 1 mL of CH<sub>2</sub>Cl<sub>2</sub>. Solvent in the vials was evaporated *in vacuo* (Genevac), leaving product residue.

#### Preparation of 5-iodoisatoic anhydride

To a red-brown solution of 2-amino-5-iodobenzoic acid (25 grams, 95 mmol) in 300  
15 mL of dioxane, 9.58 grams (32.3 mmol) of triphosgene were carefully added. The resulting slurry was refluxed for 4 hours. By that time, all starting material had disappeared by HPLC. The solid product was then filtered, washed once with ethyl ether, then dried overnight in a vacuum oven at 40 °C. The tan colored needles amounted to 22.9 grams (83 %). HPLC (MRH1 method):  $t_R = 2.15$  min. (100 %); <sup>1</sup>H  
20 NMR (400 MHz, DMSO-*d*<sub>6</sub>)  $\delta$  8.12 (s, 1 H), 8.00 (d, J = 8.6 Hz, 1 H), 6.95 (d, J = 8.5 Hz, 1 H); MS (ES) m/z (rel. intensity) 288 (M-, 100), 244 (5), 289 (5); 577 (10).

#### Preparation of 6-Chloroindoline

In a 250 mL round bottom flask, 12.4 grams of sodium cyanoborohydride (198 mmol,  
25 2 eq.) were added portion-wise over 5 minutes to a solution of 15 grams (98.9 mmol) of 6-chloroindole. After stirring for 22 hours, the mixture had become a brown solution and analysis by HPLC (MRH 1 method) revealed no starting material remaining and a mixture of two product peaks. The mixture was diluted with 100 mL of water, then made basic with ~200 mL of 6N sodium hydroxide. The desired  
30 product was extracted into 3 X 400 mL of methylene chloride. The extracts were then dried over anhydrous magnesium sulfate and evaporated *in vacuo* leaving a cloudy oil. The crude product was chromatographed over a plug of silica in 100 % methylene chloride giving a mixed fraction ( $R_f = 0.9$  and  $0.7$ ), a pure product fraction ( $R_f = 0.7$ ),

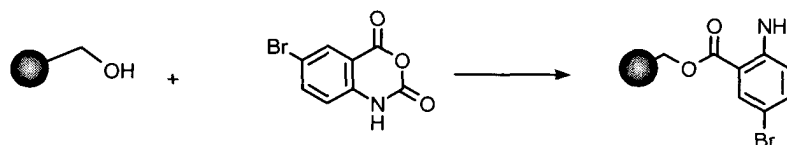
and a baseline fraction ( $R_f = 0.0 - 0.2$ ). The pure fraction was evaporated to dryness *in vacuo* to yield a clear, colorless oil weighing 10.90 grams (72 %). It was stored at 4°C and saved for future use.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ )  $\delta$  6.95 (d,  $J = 5$  Hz, 1 H), 6.46 (d,  $J = 5$  Hz, 2 H), 3.43 (t,  $J = 6$ , 2 H), 2.86 (t,  $J = 6$ , 2 H).

5

#### 10 Example 4: AMIDE DERIVATIVES

Standard procedure for attaching 5-bromoanthranilic acid to hydroxymethyl styrene resin,:

15

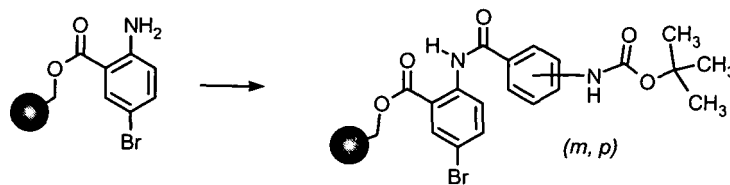


To a slurry of 24.8 g (36.7 mmol) hydroxymethyl styrene resin in 1 L DMF was added 24 g (197 mmol) 4-dimethylamino pyridine and 50 g (207 mmol) 5-bromoisatoicanhydride. The mixture was stirred at 60 °C for 18 hours and room temperature for four hours. The mixture was then filtered and the resin washed repeatedly alternating with dichloromethane and DMF (3x) then repeatedly alternating with dichloromethane and methanol (3x) followed by methanol (3x). The resin was dried over night in a vacuum oven.

25

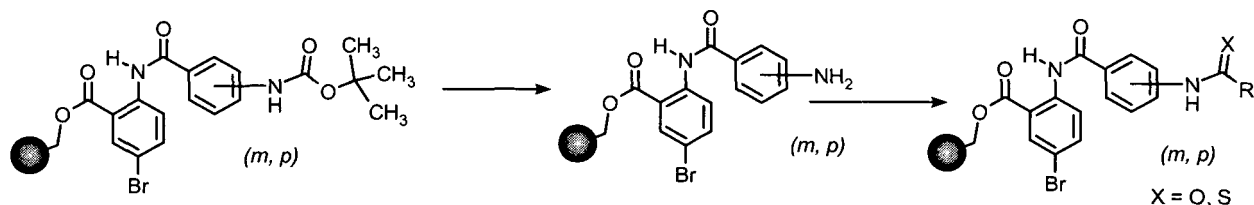
Resin 2 and 3:

Standard procedure for attaching 3 or 4- N-boc-amino benzoic acid to resin 1.



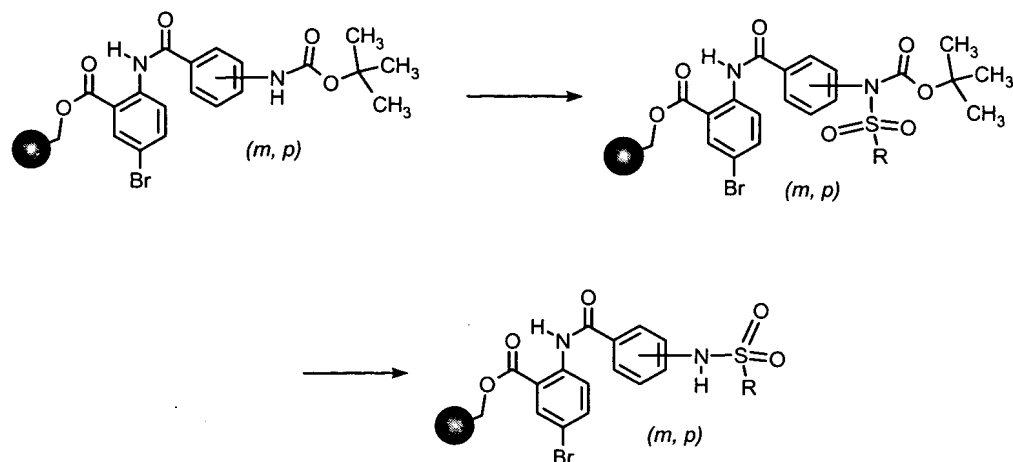
To 5.1 g (21.5 mmol) 3-N-boc-aminobenzoic acid in 200 mL of anhydrous THF was added 100  $\mu$ L DMF and 2.3 mL (25.8 mmol) oxalyl chloride in five portions over 20 minutes. After 40 minutes the mixture was concentrated in vacuo and then dissolved in 50 mL dichloromethane. This was added to a slurry of 3.79 g (4.32 mmol) resin 1 in 150 mL dichloromethane and 3.7 mL diisopropylethyl amine. The mixture was heated to reflux over night. The resin was then collected by vacuum filtration and washed repeatedly alternating with dichloromethane and methanol (4x) followed by methanol (3x) and dried in a vacuum oven. The same procedure was followed to  
 10 prepare resin 3 from 4-N-boc-aminobenzoic acid.

Standard procedure for the acylation of resins 2 and 3 with acid chlorides, isocyanates, and isothiocyanates.



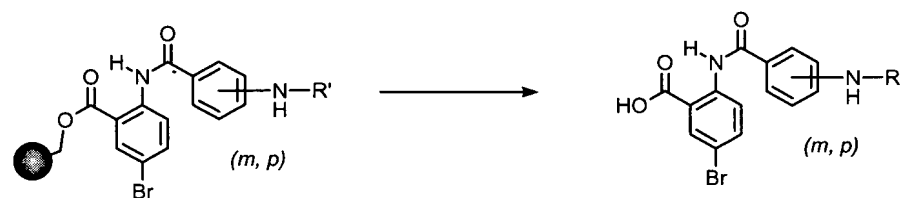
On average 55 mg (Ca. 0.055 mmol) resin was treated with 33% TFA in DCM for two hours. The resin was collected by filtration and washed repeatedly alternating with dichloromethane and methanol (4x) followed by methanol (3x) and dried in a vacuum oven. The resin is then treated with 0.6 mmol of the acylating reagent and 0.86 mmol diisopropylethyl amine in DCM and shaken over night. The resin was then collected by vacuum filtration and washed repeatedly alternating with dichloromethane and methanol (4x) followed by methanol (3x) and dried in a vacuum oven  
 20

Standard procedure for the acylation of resins 2 and 3 with sulfonyl chlorides:



On average, to 60 mg (Ca. 0.06 mmol) resin in 2 mL DCM was added 10 equivalents of a sulfonyl chloride and 174  $\mu$ L (0.6 mmol) 2-tert-butylimino-2-diethyl-amino-1,3-dimethylperhydro-1,3,2-diazaphosphorine (BEMP). After mixing overnight, the resin was collected by vacuum filtration and washed repeatedly alternating with dichloromethane and methanol (4x) followed by methanol (3x) and dried in a vacuum oven. The resin was then treated with 2 mL of 40 % TFA in DCM for one hour and then collected by vacuum filtration and washed repeatedly alternating with dichloromethane and methanol (4x) followed by methanol (3x) and dried in a vacuum oven.

Standard cleavage procedure to provide products.



The resin was treated with 1.5 mL THF and 0.5 mL 1 N sodium hydroxide over night. The mixtures were filtered and the collected filtrate was treated with 250 mg of IR-120 acidic resin for 2.5 hours. The mixtures were filtered and the filtrates concentrated to provide the following products. If initial purity was less than 80 % by HPLC those products were purified by chromatography.

Several compounds were produced by the above-described methodologies.

2-{{3-(benzoylamino)benzoyl}amino}-5-bromobenzoic acid

5-bromo-2-{{3-(2-furoylamino)benzoyl}amino}benzoic acid

5-bromo-2-({3-[(thien-2-ylacetyl)amino]benzoyl}amino)benzoic acid

- 5-bromo-2-({3-[(mesitylcarbonyl)amino]benzoyl}amino)benzoic acid  
 5-bromo-2-({4-[(mesitylcarbonyl)amino]benzoyl}amino)benzoic acid  
 2-({3-[(1,3-benzodioxol-5-ylcarbonyl)amino]benzoyl}amino)-5-bromobenzoic acid  
 5-bromo-2-({3-[(2,4-dimethoxybenzoyl)amino]benzoyl}amino)benzoic acid  
 5 5-bromo-2-[(3-{{(phenylthio)acetyl}amino}benzoyl)amino]benzoic acid  
 5-bromo-2-({3-[(methoxyacetyl)amino]benzoyl}amino)benzoic acid  
 2-({3-[(anilino)carbonyl]amino)benzoyl}amino)-5-bromobenzoic acid  
 5-bromo-2-{{3-{{(2,4-difluorophenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid  
 10 5-bromo-2-{{3-{{[(3-cyanophenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid  
 5-bromo-2-{{3-{{[(3-chlorophenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid  
 5-bromo-2-({3-{{[3-(methylthio)phenyl]amino}carbonyl}amino]benzoyl}amino)benzoic acid  
 15 2-{{3-{{[(3-acetylphenyl)amino]carbonyl}amino)benzoyl}amino}-5-bromobenzoic acid  
 5-bromo-2-({4-[(phenylsulfonyl)amino]benzoyl}amino)benzoic acid  
 5-bromo-2-{{3-{{[4-(trifluoromethoxy)phenyl]sulfonyl}amino)benzoyl}amino}benzoic acid  
 20 5-bromo-2-{{4-{{[4-(trifluoromethoxy)phenyl]sulfonyl}amino)benzoyl}amino}benzoic acid  
 5-bromo-2-[(4-{{[(3,4-dichlorophenyl)sulfonyl]amino}benzoyl}amino]benzoic acid  
 5-bromo-2-({4-[(thien-2-ylacetyl)amino]benzoyl}amino)benzoic acid  
 25 5-bromo-2-({3-[(5-nitro-2-furoyl)amino]benzoyl}amino)benzoic acid  
 5-bromo-2-({4-[(5-nitro-2-furoyl)amino]benzoyl}amino)benzoic acid  
 5-bromo-2-{{4-{{[(2,4-difluorophenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid  
 5-bromo-2-{{3-{{[(3,5-dichlorophenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid  
 30 5-bromo-2-{{3-{{[(5-chloro-2-methoxyphenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid

5-bromo-2-{{3-({[(4-phenoxyphenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid

5-bromo-2-{{4-({[(4-phenoxyphenyl)amino]carbonyl}amino)benzoyl}amino}benzoic acid

5 2-{{3-({[(4-acetylphenyl)amino]carbonyl}amino)benzoyl}amino}-5-bromobenzoic acid

5-bromo-2-{{4-({[(4-nitrophenyl)amino]carbonothioyl}amino)benzoyl}amino}benzoic acid

10 5-bromo-2-{{3-({[2-(trifluoromethyl)phenyl]amino}carbonothioyl)amino]benzoyl}amino}benzoic acid

5-bromo-2-{{3-({[(3,4,5-trimethoxyphenyl)amino]carbonothioyl}amino)benzoyl}amino}benzoic acid

5-bromo-2-{{3-({[(3-(methylthio)phenyl]amino}carbonothioyl)amino]benzoyl}amino}benzoic acid

15 2-{{3-({[(3-acetylphenyl)amino]carbonothioyl}amino)benzoyl}amino}-5-bromobenzoic acid

5-bromo-2-{{3-({[(phenylsulfonyl)amino]benzoyl}amino)benzoic acid

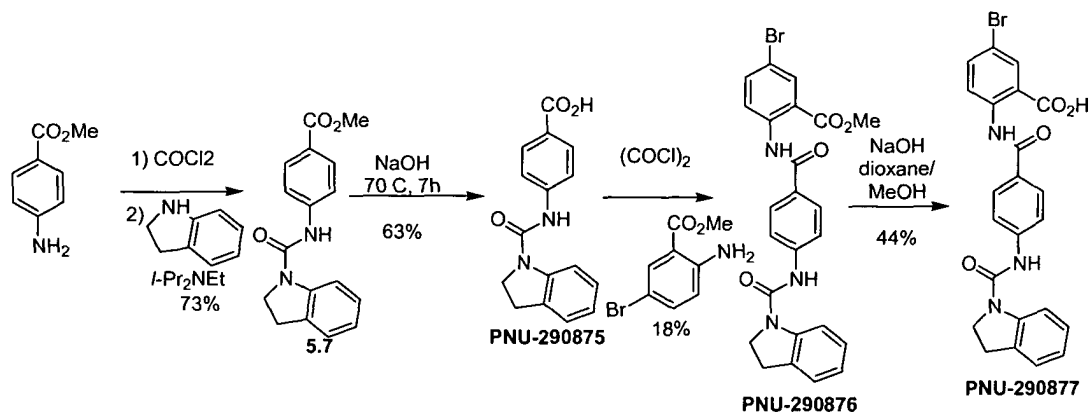
5-bromo-2-{{3-({[(3,4-dichlorophenyl)sulfonyl]amino}benzoyl)amino]benzoic acid

5-bromo-2-{{4-({[(4-methylphenyl)sulfonyl]amino}benzoyl)amino]benzoic acid

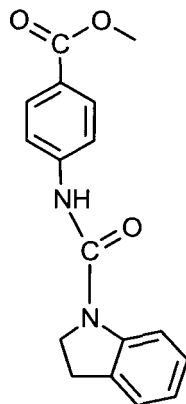
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Analogues with an alternative linkage, such as ureas, in place of the sulfonamides described in Example 1 were also synthesized.

#### Scheme 4.1



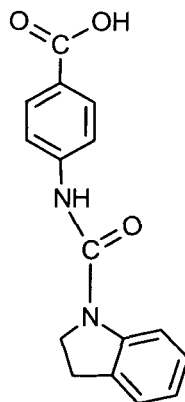
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**Methyl 4-[(2,3-dihydro-1H-indol-1-ylcarbonyl)amino]benzoate**

Methyl-4-aminobenzoate (1.00 g, 7.29 mmol) in DCM (50 mL) was slowly added to a solution of phosgene (1.93 M /toluene, 7.5 mL, 14.5 mol, 2.0 equiv) in DCM (200  
 5 mL) at 0°C, followed by the addition of diisopropylethyl amine (1.14 mL, 6.56 mmol, 0.9 equiv). The mixture was allowed to warm to rt, then stirred for 1 h, and then concentrated in vacuo to ca 5 mL. The suspension was redissolved in DCM followed by the addition of indoline (2.45 mL, 21.87 mmol, 3.0 equiv) and diisopropylethyl amine (1.14 mL, 6.56 mmol, 0.9 equiv). The resulting mixture was stirred for 2h, at  
 10 rt, then washed with 1N HCl, brine, dried (MgSO<sub>4</sub>) filtered and concentrated in vacuo. The residue was recrystallized from EtOH to afford 1.67 g of **5.7** as a white solid.

<sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 8.04-8.01 (m, 2 H), 7.90 (d, *J* = 7.9 Hz, 1 H), 7.58-7.55 (m, 2 H), 7.28-7.20 (m, 2 H), 7.01 (t, *J* = 8.2 Hz, 1 H), 6.70 (s, 1 H), 4.12 (t, *J* = 8.3 Hz, 2 H), 3.91 (s, 3 H), 3.26 (t, *J* = 8.2 Hz, 2 H).

15

**4-[(2,3-Dihydro-1H-indol-1-ylcarbonyl)amino]benzoic acid**

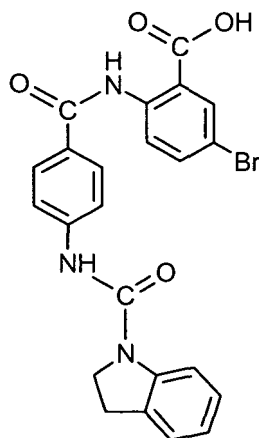


Methyl 4-[(2,3-dihydro-1H-indol-1-ylcarbonyl)amino]benzoate (1.30 g, 4.37 mmol) was placed in dioxane (50 mL) with 5 N NaOH (10 mL) and the resulting solution was heated at 70 °C for 7h. The reaction was cooled to rt, acidified, diluted with EtOAc and washed with H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>) filtered and concentrated in vacuo. The residue was recrystallized from EtOH to afford 776 mg (63%) of a white solid.

<sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.82 (s, 1 H), 7.87 (d, *J* = 8.6 Hz, 3 H), 7.71 (d, *J* = 8.7 Hz, 2 H), 7.22-7.14 (m, 2 H), 6.92 (t, *J* = 7.3 Hz, 1 H), 4.16 (t, *J* = 8.4 Hz, 2 H), 3.18 (t, *J* = 8.5 Hz, 2 H).

10

**5-Bromo-2-({4-[(2,3-dihydro-1H-indol-1-ylcarbonyl)amino]benzoyl}amino)benzoic acid, PNU-290877**



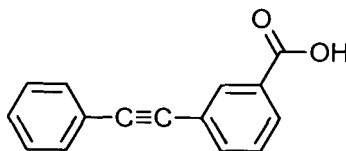
4-[(2,3-dihydro-1H-indol-1-ylcarbonyl)amino]benzoic acid (627 mg, 2.22 mmol) was dissolved in DCM (30 mL) followed by the addition of oxalyl chloride (490 µL, 5.55 mmol, 2.5 equiv) and DMF (30 µL). The mixture was stirred for 1h, then diluted with heptane (10 mL), concentrated in vacuo to dryness. The residue was redissolved in DCM (50 mL) followed by the addition of methyl-2-amino-5-bromo benzoate (510 mg, 2.2 mmol, 1.0 equiv.) and pyridine (360 µL, 4.4 mmol, 2.0 equiv.) The reaction was stirred for 3 h at rt, then washed with 1 N HCl, 1 N NaOH, H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>) filtered and concentrated in vacuo. The residue was purified by silica gel chromatography (heptane/ EtOAc 19/1, 9/1, 4/1, 1/1, 0/1) to afford 198 mg (18%) of a white solid as the methyl ester. The ester (177 mg, 0.35 mmol) was dissolved in dioxane (10 mL) followed by the addition of 5 N NaOH (5 mL). The reaction was

stirred for 3h at rt, diluted with EtOAc, washed with 1 N HCl, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was recrystallized from EtOH to afford 76 mg (44%) of a white solid.

<sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.88 (s, 1 H), 8.69 (d, *J* = 9.0 Hz, 1 H), 8.13 (d, *J* = 2.4 Hz, 1 H), 7.86-7.78 (m, 6 H), 7.21-7.14 (m, 2 H), 6.93 (t, *J* = 8.6 Hz, 1 H), 4.17 (t, *J* = 8.2 Hz, 2 H), 3.19 (t, *J* = 8.2 Hz, 1 H).

### Example 5: ALKYL DERIVATIVES

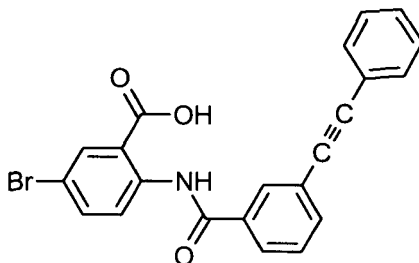
#### 10 Preparation of 3-(Phenylethynyl)benzoic acid



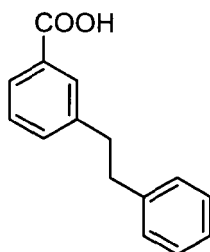
A flask containing ethyl 3-iodobenzoate (2.21g, 8.00 mmol, Lancaster), copper (I) iodide (550 mg, 2.88 mmol, Alfa), and tetrabutylammonium iodide (5.9 g, 16 mmol, Aldrich) was placed under argon. DMF (40 mL), diisopropylethylamine (4.5 mL, 26 mmol, Aldrich), and tri-*t*-butylphosphine (1.8 g of 10 wt% solution in hexane, 0.89 mmol, Strem) were added by syringe. Tris(dibenzylideneacetone)dipalladium(0)-chloroform adduct (220 mg, 0.21 mmol, Aldrich) was added as a solid under a flow or argon. The mixture was stirred for 5 minutes, and phenylacetylene (0.88 mL, 8.0 mmol, Lancaster) was added by syringe. After 40 minutes, the mixture was added to a separatory funnel with 200 mL of saturated aqueous NaHCO<sub>3</sub>. Product was extracted into 3 X 100 mL of EtOAc. The combined EtOAc was washed with 4 X 200 mL of water and then dried over MgSO<sub>4</sub>. Product was adsorbed onto silica and purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 25% - 40% CH<sub>2</sub>Cl<sub>2</sub> in heptane. The ethyl 3-(phenylethynyl)benzoate was isolated as 1.82 g of brown oil that was contaminated with tri-*t*-butylphosphine. 990 mg of this oil was dissolved in dioxane (15 mL) and treated with 1 M aqueous sodium hydroxide (6 mL), and the mixture was stirred for 3.5 hours. It was then added to a separatory funnel with 100 mL of 1 M aqueous HCl and 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. A few milliliters of THF were added to help with solubility. The organics were washed with an additional 100 mL of HCl followed by 100 mL of water and then dried over MgSO<sub>4</sub>. Solvent was removed leaving 782 mg of tan solid that was still contaminated with phosphine.

Most of this material was carried on without further purification. For the purposes of characterization, the remainder was recrystallized from ethanol/heptane yielding a white solid.

5 **5-Bromo-2-{{3-(phenylethynyl)benzoyl}amino}benzoic acid**

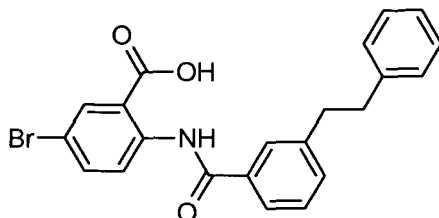


To 3-(phenylethynyl)benzoic acid (569 mg, 2.56 mmol) in  $\text{CH}_2\text{Cl}_2$  (20 mL) was added DMF (40  $\mu\text{L}$ ) and oxalyl chloride (450  $\mu\text{L}$ , 5.16 mmol). The mixture was stirred for 2.5 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (15 mL), and methyl 2-amino-5-bromobenzoate (504 mg, 2.19 mmol, Avocado) in pyridine (6 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of  $\text{CH}_2\text{Cl}_2$ . This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 50% - 60%  $\text{CH}_2\text{Cl}_2$  in heptane as eluent. Yield was 694 mg of white solid as the methyl ester. To a mixture of the methyl ester (485 mg, 1.12 mmol) in dioxane (15 mL) was added 1 M aqueous sodium hydroxide (2.2 mL). The mixture was stirred for 2.75 hours. The reaction mixture was added to a separatory funnel with 100 mL of 1 M aqueous HCl, and the product was extracted into 100 mL of  $\text{CH}_2\text{Cl}_2$ . The  $\text{CH}_2\text{Cl}_2$  was washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. It was then dried over  $\text{MgSO}_4$  and evaporated. The residue was recrystallized from hot ethanol/THF. The solids were washed with ethanol followed by heptane and then dried at 100  $^\circ\text{C}$  under vacuum yielding 295 mg of white solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  12.06 (s, 1 H), 8.60 (d,  $J$  = 9.2 Hz, 1 H), 8.12 (d,  $J$  = 2.0 Hz, 1 H), 8.10 (s, 1 H), 7.97 (d,  $J$  = 7.6 Hz, 1 H), 7.87 (dd,  $J$  = 9.2, 2.5 Hz, 1 H), 7.83 (d,  $J$  = 8.1 Hz, 1 H), 7.66 (t,  $J$  = 7.6 Hz, 1 H), 7.59-7.63 (m, 2 H), 7.45-7.48 (m, 3 H).

**Preparation of 3-(2-Phenylethyl)benzoic acid**

A mixture of 3-(phenylethynyl)benzoic acid (418 mg, 1.88 mmol) and palladium on carbon (315 mg, 10%, Aldrich) in 1:1 methanol/THF (20 mL) was stirred under 1  
 5 ATM of hydrogen overnight. The mixture was then filtered through a plug of celite and concentrated yielding 406 mg of white solid. This material was carried forward without further purification. For the purposes of characterization, a small amount of the product was recrystallized from toluene.

10

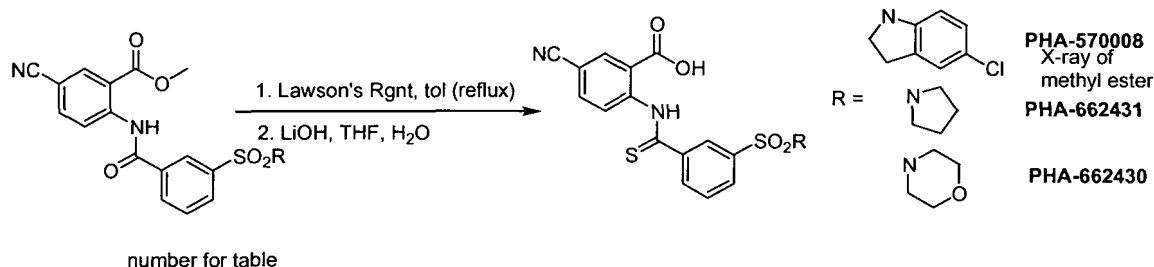
**5-Bromo-2-[[3-(2-phenylethyl)benzoyl]amino]benzoic acid**

To 3-(2-phenylethyl)benzoic acid (292 mg, 1.29 mmol) in  $\text{CH}_2\text{Cl}_2$  (20 mL) was added DMF (20  $\mu\text{L}$ ) and oxalyl chloride (225  $\mu\text{L}$ , 2.58 mmol). The mixture was stirred for  
 15 2.5 hours, and the solvent and excess oxalyl chloride were removed by rotary evaporation. The residue was dissolved in  $\text{CH}_2\text{Cl}_2$  (10 mL), and methyl 2-amino-5-bromobenzoate (248 mg, 1.08 mmol, Avocado) in pyridine (4 mL) was added. The mixture was stirred overnight and then added to a separatory funnel with 100 mL of  $\text{CH}_2\text{Cl}_2$ . This solution was washed with 2 X 100 mL of 1 M aqueous HCl and 100  
 20 mL of brine. The  $\text{CH}_2\text{Cl}_2$  was evaporated in the presence of silica gel, and the product was purified by chromatography using a Biotage Flash 40 M silica cartridge with a gradient from 50% - 100%  $\text{CH}_2\text{Cl}_2$  in heptane as eluent. Yield was 361 mg of white solid as the methyl ester. To a mixture of the methyl ester (285 mg, 0.65 mmol) in dioxane (10 mL) was added 1 M aqueous sodium hydroxide (1.0 mL). The mixture  
 25 was stirred at room temperature for 1 hour and then heated in a 50  $^\circ\text{C}$  oil bath for 15 minutes. The reaction mixture was added to a separatory funnel with 100 mL of 1 M

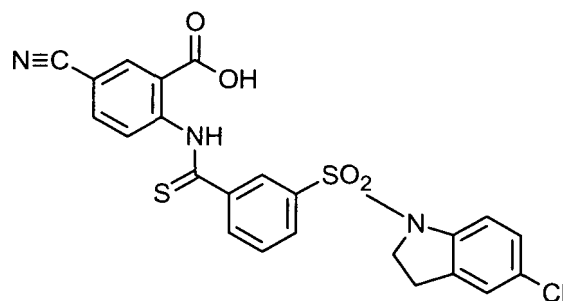
aqueous HCl, and the product was extracted into 100 mL of CH<sub>2</sub>Cl<sub>2</sub>. The CH<sub>2</sub>Cl<sub>2</sub> was washed with an additional 100 mL of 1 M aqueous HCl followed by 100 mL of water. It was then dried over MgSO<sub>4</sub> and evaporated. The residue was recrystallized from hot ethanol. The solids were washed with heptane and then dried at 100 °C under vacuum yielding 88 mg of white solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 12.10 (s, 1 H), 8.68 (d, *J* = 9.1 Hz, 1 H), 8.12 (d, *J* = 2.5 Hz, 1 H), 7.83-7.87 (m, 2 H), 7.75-7.78 (m, 1 H), 7.46-7.51 (m, 2 H), 7.16-7.31 (m, 5 H), 2.91-3.02 (m, 4 H).

### Example 6:

#### Thioamide linkers.



#### 2-[(3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl)carbonothioyl]amino]-5-cyanobenzoic acid

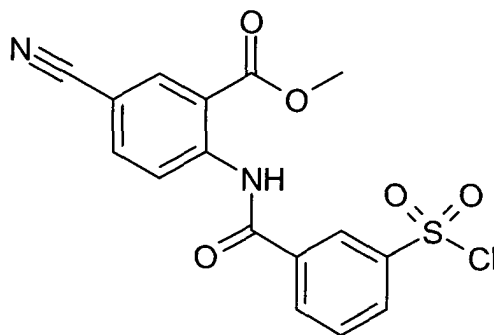


General procedure A: Methyl 2-[(3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl)carbonyl] amino]-5-cyanobenzoate (989 mg, 1.99 mmol) and Lawesson's reagent (4.5 g, 11.1 mmol) were combined in a flask equipped with a reflux condensor. The flask was evacuated and purged with N<sub>2</sub> several times. Tol (30 mL) was added and the reaction was refluxed overnight. The reaction was cooled to rt and filtered to remove excess Lawesson's reagent. The filtrate was absorbed in SiO<sub>2</sub>

and the product was purified by silica gel chromatography using Hept/EtOAc (19:1, 9:1, 3:17, 4:1). The product was triturated with MeOH to afford 670 mg (66%) of an orange solid as the methyl ester. <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 12.40 (s, 1 H), 8.35 (d, *J* = 2 Hz, 1 H), 8.29 (s, 1 H), 8.19 (dd, *J* = 8, 2 Hz, 1 H), 8.14 (d, *J* = 8 Hz, 1 H), 8.06 (d, *J* = 8 Hz, 1 H), 7.98 (d, *J* = 8 Hz, 1 H), 7.73 (t, *J* = 8 Hz, 1 H), 7.49 (d, *J* = 9 Hz, 1 H), 7.30-7.25 (m, 2 H), 4.02 (t, *J* = 8 Hz, 2 H), 3.79 (s, 3 H), 2.97 (t, *J* = 8 Hz, 2 H).

General procedure B: to a solution of the methyl ester (300 mg, 0.605 mmol) dissolved in THF (7 mL) and H<sub>2</sub>O (1.5 mL) was added LiOH-H<sub>2</sub>O (450 mg, 10.7 mmol) and the reaction was heated to 45°C for 6 hr. The solution was diluted with MTBE, washed with 2 N HCl and brine, dried (MgSO<sub>4</sub>), concentrated, and triturated with MeOH to afford 252 mg (84%) of an orange solid. <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 8.62 (d, *J* = 8 Hz, 1 H), 8.36 (dd, *J* = 12, 2 Hz, 1 H), 8.18 (d, *J* = 8 Hz, 1 H), 8.12 (dd, *J* = 8, 2 Hz, 1 H), 7.95 (d, *J* = 8 Hz, 1 H), 7.71 (t, *J* = 8 Hz, 1 H), 7.48 (d, *J* = 9 Hz, 1 H), 7.27-7.25 (m, 2 H), 4.02 (t, *J* = 8 Hz, 2 H), 2.96 (t, *J* = 8 Hz, 2 H).

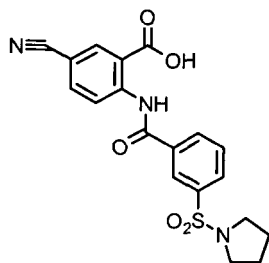
**Methyl 2-{[3-(chlorosulfonyl)benzoyl]amino}-5-cyanobenzoate**



To a suspension of 3-(chlorosulfonyl)benzoic acid (10.8 g, 49.0 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (105 mL) and three drops of DMF was added oxalyl chloride (12.5 mL) and the reaction was stirred at rt overnight. The solution was concentrated *in vacuo*, diluted with CH<sub>2</sub>Cl<sub>2</sub> (100 mL), and the solution was divided into two reactions. A 50 mL (24.5 mmol) aliquot of the acid chloride was added to a solution of PHA-522499 (4.49 g, 25.5 mmol) dissolved in CH<sub>2</sub>Cl<sub>2</sub> (50 mL) and pyridine (3.0 mL) and stirred at rt overnight. The solution was diluted with MTBE, washed with 2 N HCl and brine, concentrated, triturated with MTBE to afford 7.91 g (85%) of methyl 2-{[3-(chlorosulfonyl)benzoyl]amino}-5-cyanobenzoate as a tan solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 11.73 (s, 1 H), 8.67 (d, *J* = 9 Hz, 1 H), 8.37 (d, *J* = 2 Hz, 1 H), 8.25 (s, 1

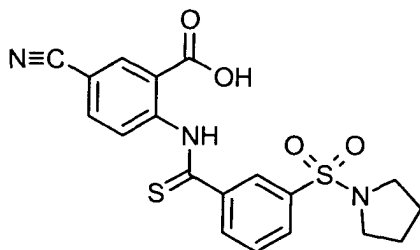
H), 8.12 (dd,  $J = 9, 2$  Hz, 1 H), 7.92 (d,  $J = 8$  Hz, 1 H), 7.88 (d,  $J = 8$  Hz, 1 H), 7.60 (t,  $J = 8$  Hz, 1 H), 3.93 (s, 3 H).

5 **5-cyano-2-{{3-(pyrrolidin-1-ylsulfonyl)benzoyl}amino}benzoic acid**



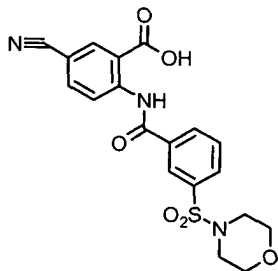
General procedure C: To a solution of methyl 2-{{3-(chlorosulfonyl)benzoyl}amino}-5-cyanobenzoate (1.863 g, 4.92 mmol) dissolved in  $\text{CH}_2\text{Cl}_2$  (40 mL) was added  
 10 pyrrolidine (1.5 mL, 18.0 mmol) and stirred at rt for 3 hr. The reaction was diluted with MTBE, washed with 2 N HCl and brine, concentrated, and triturated with MeOH to afford 1.70 g (84%) of methyl 5-cyano-2-{{3-(pyrrolidin-1-ylsulfonyl)benzoyl}amino}benzoate as a tan solid.  $^1\text{H}$  NMR (300 MHz,  $\text{DMSO}-d_6$ )  $\delta$  11.75 (s, 1 H), 8.61 (d,  $J = 9$  Hz, 1 H), 8.38 (d,  $J = 2$  Hz, 1 H), 8.33 (s, 1 H), 8.25 (d,  $J = 8$  Hz, 1 H), 8.14 (dd,  $J = 9, 2$  Hz, 1 H), 8.10 (d,  $J = 8$  Hz, 1 H), 7.90 (t,  $J = 8$  Hz, 1 H), 3.91 (s, 3 H), 3.24-3.19 (m, 4 H), 1.71-1.66 (m, 4 H). Methyl 2-{{3-(chlorosulfonyl)benzoyl}amino}-5-cyanobenzoate (378 mg, 1.0 mmol) was dissolved in 15 mL of  $\text{CHCl}_3$ . Pyrrolidine (145 mg, 2.0 mmol) and  $\text{Et}_3\text{N}$  (1 mL) were then added and the reaction stirred at room temperature for 12 hr. The mixture was poured  
 20 into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over  $\text{Na}_2\text{SO}_4$  and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography, providing 297 mg (72%) of the desired methyl ester. The ester was treated with LiOH in 1:1:1 THF/MeOH/ $\text{H}_2\text{O}$  for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over  $\text{Na}_2\text{SO}_4$  and then concentrated *in vacuo*. The title compound (249 mg, 87%) was  
 25 obtained as a white solid after recrystallization from MeOH.  $^1\text{H}$  NMR (300 MHz, DMSO) 1.67 (m, 4H), 3.20 (m, 4H), 7.88 (t, 1H), 8.09-8.14 (m, 2H), 8.26 (d, 1H), 8.33 (s, 1H), 8.42 (d, 1H), 8.83 (d, 1H), 12.56 (s, 1H)

**5-Cyano-2-([3-(pyrrolidin-1-ylsulfonyl)phenyl]carbonothioyl)amino)benzoic acid**



Prepared according to general procedure A: Methyl 5-cyano-2-([3-(pyrrolidin-1-ylsulfonyl)benzoyl]amino)benzoate (1.12 g, 2.70 mmol) and Lawesson's reagent (5.5 g, 13.6 mmol) afforded 450 mg of a mixture of the methyl ester and Lawesson's reagent after purifying by silica gel chromatography twice. The crude material was hydrolyzed according to general method B to afford 253 mg (29%) over two steps of an orange solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 9.80 (d, *J* = 9 Hz, 1 H), 8.42 (d, *J* = 2 Hz, 1 H), 8.33 (s, 1 H), 8.23 (d, *J* = 8 Hz, 1 H), 7.97-7.91 (m, 2 H), 7.75 (t, *J* = 7 Hz, 1 H), 3.23-3.19 (m, 4 H), 1.71-1.65 (m, 4 H).

**5-cyano-2-([3-(morpholin-4-ylsulfonyl)benzoyl]amino)benzoic acid**

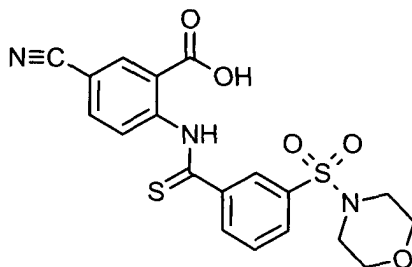


Methyl 2-([3-(chlorosulfonyl)benzoyl]amino)-5-cyanobenzoate (378 mg, 1.0 mmol) was dissolved in 15 mL of CHCl<sub>3</sub>. Morpholine (156 mg, 2.0 mmol) and Et<sub>3</sub>N (1 mL) were then added and the reaction stirred at room temperature for 12 hr. The mixture was poured into 1 M HCl (20 mL) and extracted with EtOAc (3 x 20 mL). The combined organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. The resulting residue was purified by silica gel chromatography, providing 373 mg (87%) of the desired methyl ester. The ester was treated with LiOH in 1:1:1 THF/MeOH/H<sub>2</sub>O for 12 hrs followed by acidification and extraction with EtOAc. The organic solution was dried over Na<sub>2</sub>SO<sub>4</sub> and then concentrated *in vacuo*. The title compound (298 mg, 82%) was obtained as a white solid after recrystallization from



MeOH.  $^1\text{H}$  NMR (400 MHz, DMSO) 2.94 (m, 4H), 3.65 (m, 4H), 7.96 (t, 1H), 8.03 (d, 1H), 8.13 (dd, 1H), 8.27-8.31 (m, 2H), 8.42 (d, 1H), 8.82 (d, 1H), 12.55 (s, 1H)

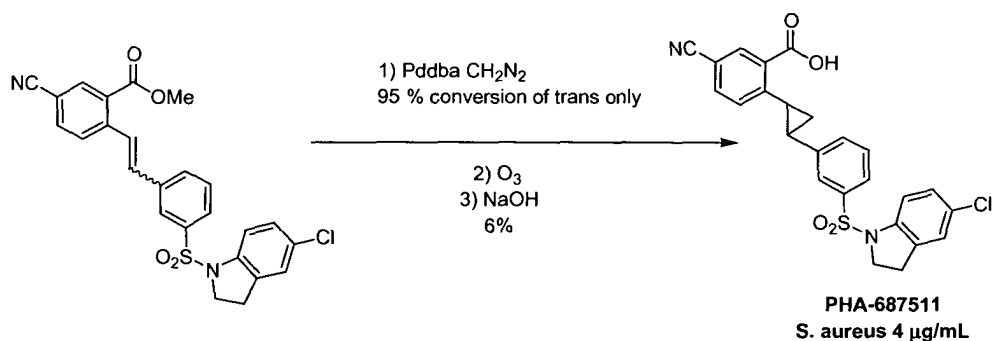
**5-Cyano-2-([3-(morpholin-4-ylsulfonyl)phenyl]carbonothioyl)amino)benzoic acid**



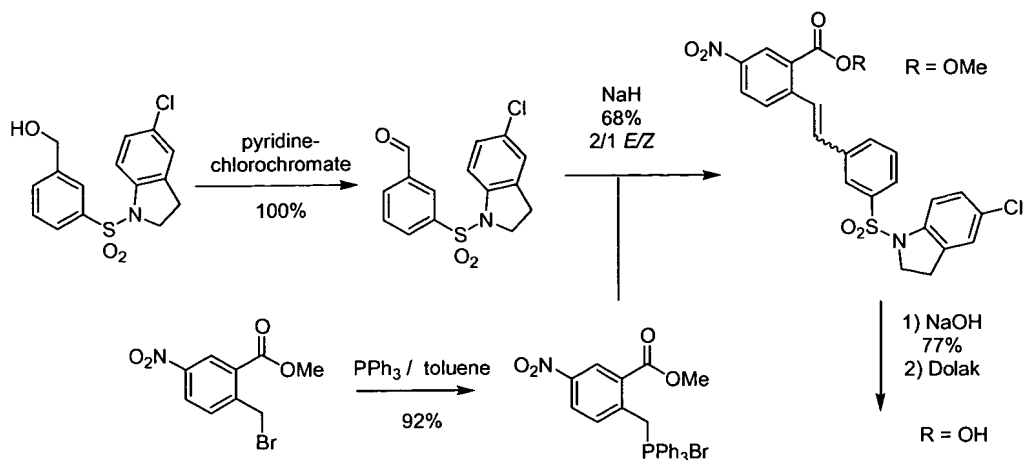
Prepared according to general method A and B: Methyl 5-cyano-2-([3-(morpholin-4-ylsulfonyl)benzoyl]amino)benzoate (1.02 g, 2.38 mmol) and Lawesson's reagent (4.78 g, 11.8 mmol) afforded 532 g (50%) of the ester, 35527-bdw-118 as an orange solid. The ester (495 mg, 1.09 mmol) was hydrolyzed by general procedure B to afford 87 mg (20%) of an orange solid.  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ )  $\delta$  9.72 (d,  $J$  = 8 Hz, 1 H), 8.41 (d,  $J$  = 2 Hz, 1 H), 8.27-8.25 (m, 2 H), 7.95 (dd,  $J$  = 9, 6 Hz, 1 H), 7.90 (d,  $J$  = 9 Hz, 1 H), 7.79 (t,  $J$  = 6 Hz, 1 H).

**Example 7: X-Y Derivatives**

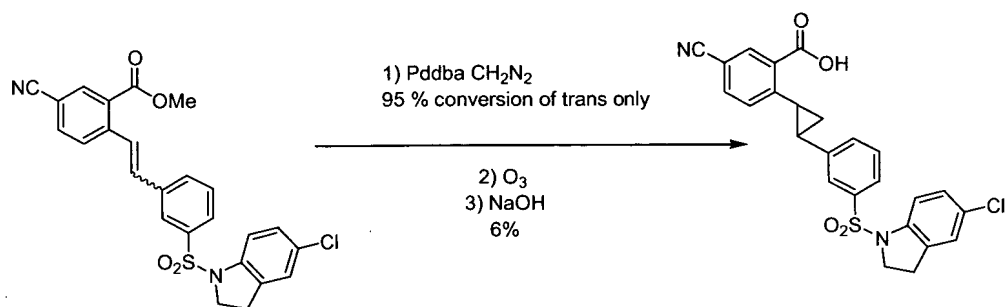
Scheme 7.1



**Scheme 7.2**

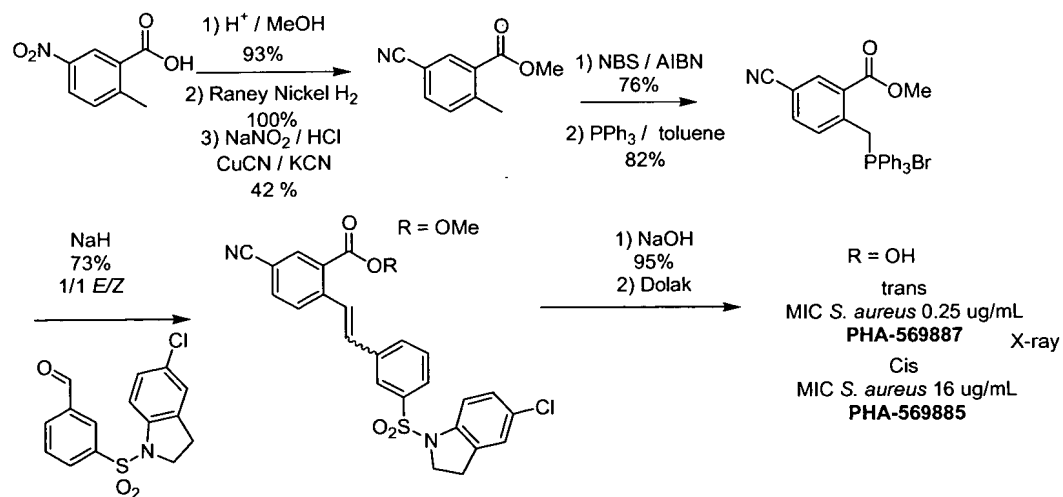


Scheme 7.3

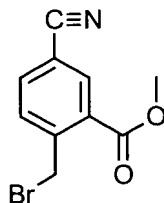


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Scheme 7.4

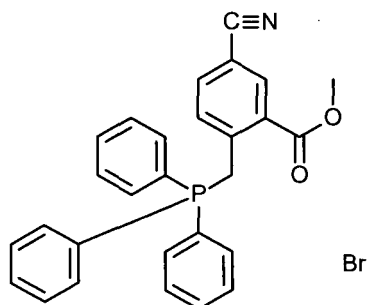


## Methyl 2-(bromomethyl)-5-cyanobenzoate



Methyl 5-cyanobenzoate (4.50 g, 25.6 mmol), NBS (5.03 g, 28.25 mmol) and AIBN (150 mg) were dissolved in dichloroethane (160 mL). The mixture was irradiated with a photolamp for 2h. The mixture was cooled to rt and concentrated in vacuo. The residue was purified by silica gel chromatography (DCM/heptane 1/9, 1/4, 1/1, 1/0) to afford 4.79 g (73%) of methyl 2-(bromomethyl)-5-cyanobenzoate. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 8.29 (d, *J* = 1.7 Hz, 1 H), 7.79 (dd, *J* = 8.0, 1.7 Hz, 1 H), 7.63 (d, *J* = 8.0 Hz, 1 H), 4.97 (s, 2 H), 4.00 (s, 3 H).

### Methyl 2-{{bromo(triphenyl)phosphoranyl}methyl}-5-cyanobenzoate

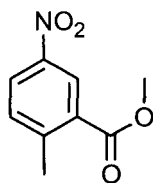


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Methyl 2-(bromomethyl)-5-cyanobenzoate (2.80 g, 10.9 mol) was added to a solution of triphenylphosphine (2.87 g, 10.9 mmol) in toluene (50 mL). The resulting mixture was heated at reflux for 3h, cooled to rt, the precipitate was isolated by filtration, washed with pentane to afford 4.64 g (82%) of methyl 2-{{bromo(triphenyl)phosphoranyl}methyl}-5-cyanobenzoate as a white solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.22 (s, 1 H), 8.08 (d, *J* = 7.9 Hz, 1 H), 8.79-7.51 (m, 16 H), 5.63 (d, *J* = 16.2 Hz, 2 H), 3.48 (s, 3 H).

15

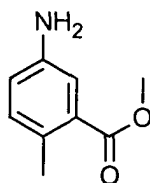
### Methyl 2-methyl-5-nitrobenzoate



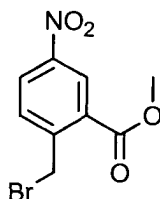
20

2-Methyl-5-nitrobenzoate (5.0 g, 27.6 mmol) was dissolved in MeOH (0.4 L) followed by the addition of H<sub>2</sub>SO<sub>4</sub> (7 mL). The mixture was heated at reflux for 36 h, then cooled to rt and concentrated to ca 100 mL. The solution was diluted with MTBE neutralized with 6N NaOH, washed with 1N NaOH, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo to afford 4.72 g (87%) of methyl 2-methyl-5-nitrobenzoate as a white solid.

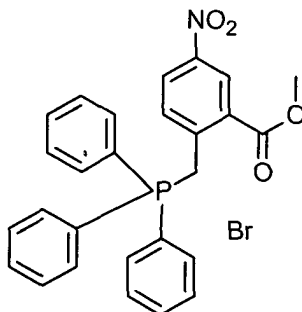
25

**Methyl 5-amino-2-methylbenzoate**

Methyl 2-methyl-5-nitrobenzoate (5.0 g, 25.6 mmol) was dissolved in EtOH with  
 5 Raney nickel under a 35 psi atmosphere of H<sub>2</sub>. The reaction was stirred for 20 h, then  
 filtered through Celite washed with MeOH and concentrated in vacuo to afford 4.2 g  
 (100%) of methyl 5-amino-2-methylbenzoate.

**Methyl 2-(bromomethyl)-5-nitrobenzoate**

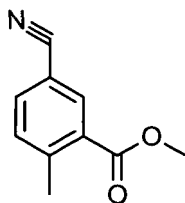
10 Methyl 2-methyl-5-nitrobenzoate (2.0 g, 10.2 mmol) NBS (2.73 g, 15.3 mmol) and  
 AIBN (50 mg) were dissolved in dichloroethane (100 mL). The mixture was irradiated  
 with a photolamp for 3h. The mixture was cooled to rt and concentrated in vacuo.  
 The residue was purified by silica gel chromatography (heptane/EtOAc 1/0, 19/1, 9/1)  
 15 to afford 2.40 g (85%) of methyl 2-(bromomethyl)-5-nitrobenzoate.

**Methyl 2-[[bromo(triphenyl)phosphoranyl]methyl]-5-nitrobenzoate**

Methyl 2-(bromomethyl)-5-nitrobenzoate (666 mg, 2.43 mmol) was added to a  
 20 solution of triphenylphosphine (640 mg, 2.4 mmol) in toluene (20 mL). The resulting  
 mixture was heated at reflux for 3h, cooled to rt, the precipitate was isolated by

filtration, washed with pentane to afford 1.2 g (92%) of methyl 2-  
 {[bromo(triphenyl)phosphoranyl]methyl}-5-nitrobenzoate as a white solid.

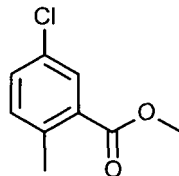
### Methyl 5-cyano-2-methylbenzoate



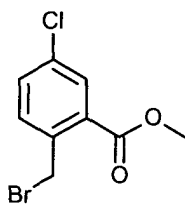
5

Methyl 5-amino-2-methylbenzoate (4.2 g, 25.4 mmol) was dissolved in MeOH/H<sub>2</sub>O (20 mL/46 mL) was cooled with icebath followed by the addition of HCl (54 mL), NaNO<sub>2</sub> (2.63 g, 38.1 mmol, in H<sub>2</sub>O 60 mL). The mixture was stirred for ½ h, then neutralized with solid NaHCO<sub>3</sub>, extensive gasevolution. Then a cold mixture of KCN  
 10 (2.48 g, 38 mmol) and CuCN (2.9 g, 33 mmol) in a H<sub>2</sub>O (40 ml)/ EtOAc (80 mL) was added. The reaction was stirred for ½ h, then filtered through Celite, extracted with EtOAc then washed with H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was purified by silica gel chromatography (heptane/DCM 19/1, 9/1, 1/1, 1/0) to afford 1.89 g (42%) of a white solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ  
 15 8.23 (d, *J* = 1.7 Hz, 1 H), 7.68 (dd, *J* = 1.8, 7.9 Hz, 1 H), 7.38 (d, *J* = 7.9 Hz, 1 H), 3.94 (s, 3 H).

### Methyl 5-chloro-2-methylbenzoate

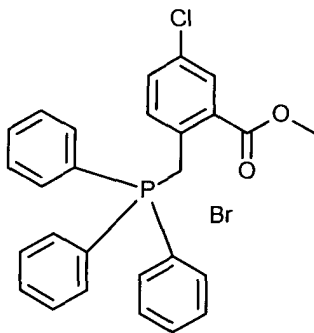


20 Methyl 5-chloro-2-methylbenzoate (25 g, 147 mmol) was dissolved in MeOH (0.6 L) followed by the addition of H<sub>2</sub>SO<sub>4</sub> (50 mL). The mixture was heated at reflux for 12 h, then cooled to rt and concentrated to ca 200 mL. The solution was diluted with MTBE, washed with H<sub>2</sub>O, 1N NaOH, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo to afford 24.9 g (92%) of methyl 5-chloro-2-methyl-benzoate as a white  
 25 solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 7.91 (d, *J* = 2.3 Hz, 1 H), 7.38 (dd, *J* = 2.3, 8.1 Hz, 1 H), 7.19 (d, *J* = 8.2 Hz, 1 H), 3.91 (s, 3 H).

**Methyl 2-(bromomethyl)-5-chlorobenzoate**

Methyl 5-chloro-2-methyl benzoate (10.0 g, 54 mmol) NBS (10.6 g, 59.5 mmol) and AIBN (200 mg) were dissolved in dichloroethane (300 mL). The mixture was irradiated with a photolamp for 2h. The mixture was cooled to rt and concentrated in vacuo. The residue was purified by silica gel chromatography (heptane/DCM 9/1, 4/1, 1/1) to afford 11.8 g (83%) of methyl 2-(bromomethyl)-5-chlorobenzoate. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 7.98 (d, *J* = 2.1 Hz, 1 H), 7.49 (dd, *J* = 2.2, 8.2 Hz, 1 H), 7.43 (d, *J* = 8.2 Hz, 1 H), 4.93 (s, 2 H), 3.97 (s, 3 H).

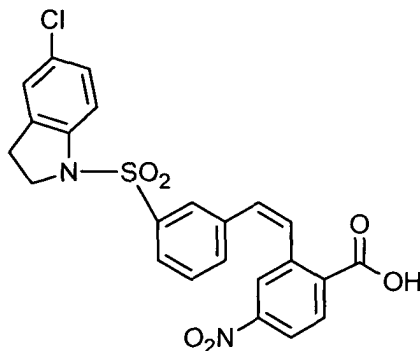
10

**Methyl 2-[[bromo(triphenyl)phosphoranyl]methyl]-5-chlorobenzoate**

Methyl 2-(bromomethyl)-5-chlorobenzoate (11.8 g, 44.6 mmol) was added to a solution of triphenylphosphine (11.6 g, 44.6 mmol) in toluene (400 mL). The resulting mixture was heated at reflux for 3h, cooled to rt, the precipitate was isolated by filtration, washed with pentane to afford 18.7 g (80%) of methyl 2-[[bromo(triphenyl) phosphoranyl] methyl]-5-chlorobenzoate as a white solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 7.85-7.68 (m, 5 H), 7.63-7.57 (m, 12 H), 7.38-7.28 (m, 1 H), 5.88 (d, *J* = 15.0 Hz, 2 H), 3.43 (s, 3 H).

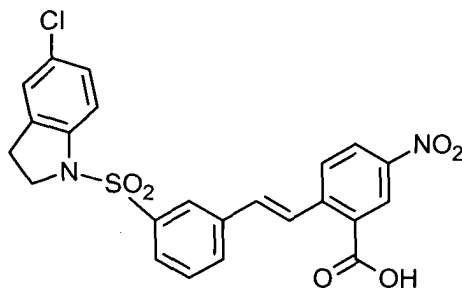
20

**2-((Z)-2-{3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}ethenyl)-4-nitrobenzoic acid**



Methyl 2-[[bromo(triphenyl)phosphoranyl]methyl]-5-nitrobenzoate (1.20 g, 2.24 mmol) was added to DMSO (30 mL) followed by NaH (100 mg, 2.4 mmol), gas evolution was observed, and the resulting mixture was heated at 60 °C for 2h. Then  
 5 3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzaldehyde (800 mg, 2.5 mmol) in toluene (50 mL) was added the reaction was stirred at rt for 2h, then at 60 °C for 2h. The mixture was diluted with MTBE, washed with H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was purified by silica gel chromatography (DCM/MeOH 1/0, 19/1) to afford 760 mg (68%) of a Z/E mixture  
 10 (4/1). The solid was dissolved in THF/MeOH (2/1, 60 mL) and 6N NaOH (6 mL) was added. The mixture was stirred at rt for 1 h, then diluted with MTBE, washed with 1N HCl, H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was purified by silica gel chromatography (DCM/MeOH 1/0, 19/1) to afford 574 mg (77%). This was recrystallized from MeOH. The mother liquid was  
 15 recrystallized several time to afford 182 mg. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 8.66 (d, *J* = 2.4 Hz, 1 H), 8.14-8.12 (m, 1 H), 7.63-7.57 (m, 1 H), 7.49-7.18 (m, 8 H), 6.84 (d, *J* = 12.3 Hz, 1 H), 3.65 (t, *J* = 8.4 Hz, 2 H), 2.86 (t, *J* = 8.4 Hz, 2 H).

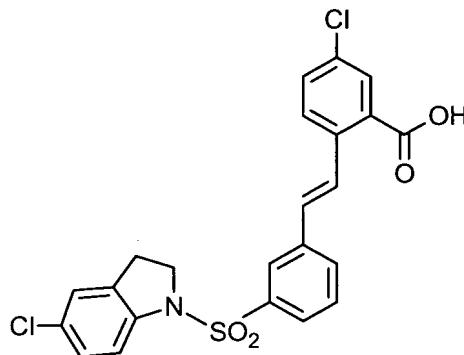
20 **2-((E)-2-[[3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl]ethenyl]-4-nitrobenzoic acid,**



<sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 8.56 (d, *J* = 2.4 Hz, 1 H), 8.33-8.31 (m, 1 H), 8.18 (d, *J* = 16.5 Hz, 1 H), 8.08-8.03 (m, 2 H), 7.92 (d, *J* = 7.8 Hz, 1 H), 7.75-7.73 (m, 1 H), 7.62 (t, *J* = 7.8 Hz, 1 H), 7.51-7.47 (m, 2 H), 7.27-7.23 (m, 2 H), 3.98 (t, *J* = 8.4 Hz, 2 H), 2.94 (t, *J* = 8.4 Hz, 2 H).

5

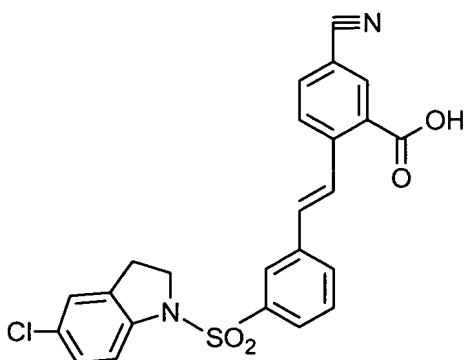
**5-Chloro-2-((E)-2-{3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}ethenyl)benzoic acid**



Methyl 2-[[bromo(triphenyl)phosphoranyl]methyl]-5-chlorobenzoate (392 mg, 0.74 mmol) was added to THF (10 mL) in icebath, followed by LiCl (260 mg, 6.2 mmol), and *n*-BuLi (300 μL, 0.74 mmol). The reaction was stirred at rt for 10 min, then 3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzaldehyde (200 mg, 0.6 mmol) was added and the reaction was stirred at rt for 2h. The mixture was diluted with MTBE, washed with H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was purified by silica gel plug (DCM) to afford 271 mg of a Z/E mixture. The solid was dissolved in toluene (10 mL) followed by the addition of thiophenol (32 μL, 0.28 mmol) and AIBN (14 mg, 0.08 mmol). The reaction was heated at reflux for 12 h, then concentrated in vacuo. The residue was dissolved in THF (60 mL) and 6N NaOH (5 mL) was added. The mixture was stirred at 100 °C for 4 h, then diluted with MTBE, washed with 1N HCl, H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was recrystallized from MeOH to afford 123 mg. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 7.97-7.85 (m, 5 H), 7.70-7.60 (m, 3 H), 7.48 (d, *J* = 8.2 Hz, 1 H), 7.33-7.24 (m, 3 H), 3.97 (t, *J* = 8.4 Hz, 2 H), 2.93 (t, *J* = 8.4 Hz, 2 H).

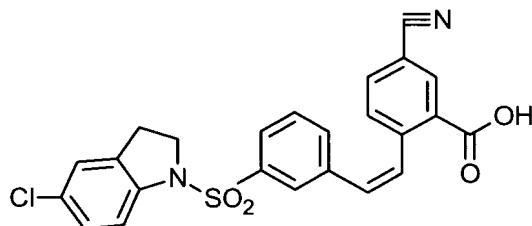
**5-Cyano-2-((E)-2-{3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}ethenyl)benzoic acid**





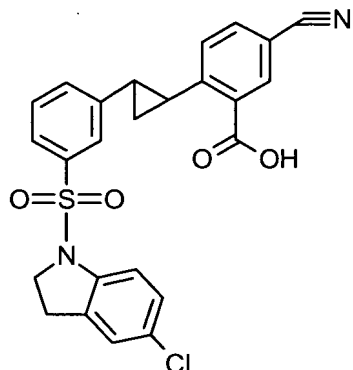
Methyl 2-[[bromo(triphenyl)phosphoranyl]methyl]-5-cyanobenzoate (1.36 g, 2.6 mmol) was added to DMSO (20 mL) followed by NaH (105 mg, 2.6 mmol), gas evolution was observed, and the resulting mixture was heated at 60 °C for 2h. Then  
 5 3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzaldehyde (564 mg, 1.7 mmol) in toluene (50 mL) was added the reaction was stirred at rt for 1h, then at 60 °C for 1h. The mixture was diluted with MTBE, washed with H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo. The residue was purified by silica gel chromatography (DCM/heptane 1/1, 1/0) to afford 616 mg (73%) of a Z/E mixture.  
 10 The solid was dissolved in THF (60 mL) and 1N NaOH (10 mL) was added. The mixture was stirred at rt for 12 h, then diluted with MTBE, washed with 1N HCl, H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo to afford 567 mg (95%). This was purified by preparative reverse phase HPLC to afford 144 mg of pure (E) and 99 mg of (Z). <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.24 (s, 1 H), 8.05-7.89 (m, 5 H), 7.76-7.73 (m, 1 H), 7.63 (t, *J* = 7.7 Hz, 1 H), 7.49-7.44 (m, 2 H), 7.27-7.24 (m, 2 H), 3.98 (t, *J* = 8.5 Hz, 2 H), 2.93 (t, *J* = 8.5 Hz, 2 H).  
 15

**5-Cyano-2-((Z)-2-{3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}ethenyl)benzoic acid**



20 <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 8.33 (d, *J* = 1.7 Hz, 1 H), 7.84-7.81 (m, 1 H), 7.59-7.57 (m, 1 H), 7.47-7.12 (m, 8 H), 6.82 (d, *J* = 12.2 Hz, 1 H), 3.66 (t, *J* = 8.5 Hz, 2 H), 2.88 (t, *J* = 8.3 Hz, 2 H).

**2-(2-{3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}cyclopropyl)-5-cyanobenzoic acid**



5 Diazomethane solution (400 ml, from 36 g Dizald, for procedure see Denmark, S. E.; Stavenger, R. A.; Faucher, A-M.; Edwards, J. P. *J. Org. Chem.* **1997**, 62, 3375) was added to a solution of methyl 5-cyano-2-(2-{3-[(5-chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl] phenyl}ethenyl)benzoate (850 mg, 1.7 mmol) and Pdca (100 mg) in DCM (150 mL). Extensive gas evolution was observed, the resulting mixture was stirred for

10 12 h, then HOAc (5 mL) was added, filtered through Celite, washed with 1N NaOH, brine, dried (MgSO<sub>4</sub>), filtered and concentrated in vacuo to afford 982 mg of a solid. The residue in DCM (100 mL) was cooled with icebath and O<sub>3</sub> was bubbled through for 30 min. Then NaBH<sub>4</sub> (500 mg) was added and the mixture was stirred for 30 min at rt. The mixture was passed through a silica plug and concentrated in vacuo. The

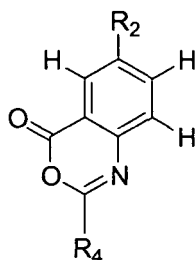
15 residue was purified by silica gel chromatography (heptane/DCM 9/1, 4/1, 1/1, 0/1) to afford 124 mg of the desired cyclopropane. The solid was dissolved in THF (25 mL) and 6N NaOH (5 mL) was added, the resulting mixture was stirred for 16h at rt, then diluted with MTBE, washed with 1N HCl, H<sub>2</sub>O, brine, dried (MgSO<sub>4</sub>) filtered and concentrated in vacuo. The residue was purified by silica gel chromatography

20 (DCM/MeOH 1/0, 19/1, 9/1, 4/1) to afford 51 mg (6%). <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>) δ 8.27 (d, *J* = 1.6 Hz, 1 H), 7.79 (d, *J* = 8.1 Hz, 1 H), 7.60-7.57 (m, 3 H), 7.40 (d, *J* = 5.3 Hz, 2 H), 7.25 (d, *J* = 8.1 Hz, 1 H), 7.18-7.16 (m, 1 H), 7.04 (s, 1 H), 3.95 (t, *J* = 8.3 Hz, 2 H), 3.18-3.13 (m, 1 H), 2.87 (t, *J* = 8.3 Hz, 2 H), 2.24-2.19 (m, 1 H), 1.65-1.60 (m, 1 H), 1.54-1.49 (m, 1 H).

25

**Example 8:**

In other embodiments, the invention includes benzoxazine derivatives of the formula



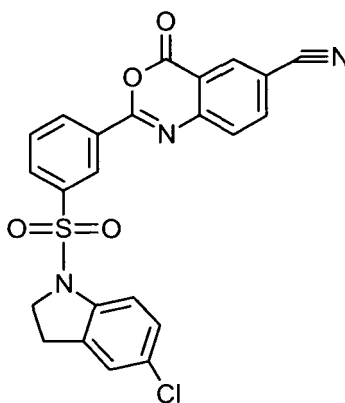
5 wherein

$R_2$  is an electron withdrawing group; and

$R_4$  is an optionally substituted aryl, provided that the aryl is not simultaneously substituted with a sulfonamide and a urea or thiourea, and further provided that the aryl is not solely substituted at the ortho-position relative to Y.

10

**2-{3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]phenyl}-4-oxo-4H-3,1-benzoxazine-6-carbonitrile**



15 2-({3-[(5-Chloro-2,3-dihydro-1H-indol-1-yl)sulfonyl]benzoyl}amino)-5-cyanobenzoic acid (PHA-524523, 884 m, 1.84 mmol) was dissolved in anhydrous THF (30 mL) and  $\text{Et}_3\text{N}$  (0.563 mL, 4.04 mmol) under  $\text{N}_2$ . Addition of ethyl chloroformate (0.193 mL, 2.02 mmol, Aldrich) to the yellow solution produced a white precipitate, which was stirred overnight at RT. The solvent was evaporated and the resultant residue

20 suspended in  $\text{CH}_2\text{Cl}_2$  (100 mL). The organic layer was washed 2x with 1.0M HCl, 1x with water and 1x with brine (100 mL each). The crude product was purified on a Biotage Flash 40M (90 g) silica cartridge using a step gradient of 0-1%  $\text{CH}_3\text{OH}$  in  $\text{CH}_2\text{Cl}_2$ . After evaporation the resultant solid was dried under vacuum at 100 °C to

afford 280 mg (33%) of white solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  8.65 (d,  $J$  = 1.9 Hz, 1 H), 8.52 (s, 1 H), 8.47 (d,  $J$  = 8.0 Hz, 1 H), 8.36 (dd,  $J$  = 8.4, 1.9 Hz, 1 H), 8.11 (d,  $J$  = 8.4 Hz, 1 H), 7.92 (d,  $J$  = 8.4 Hz, 1 H), 7.85 (t,  $J$  = 7.9 Hz, 1 H), 7.53 (d,  $J$  = 8.6 Hz, 1 H), 7.30 (d,  $J$  = 8.6 Hz, 1 H), 7.26 (s, 1 H), 3.99 (t,  $J$  = 8.4 Hz, 2 H), 2.94 (t,  $J$  = 8.4 Hz, 2 H).

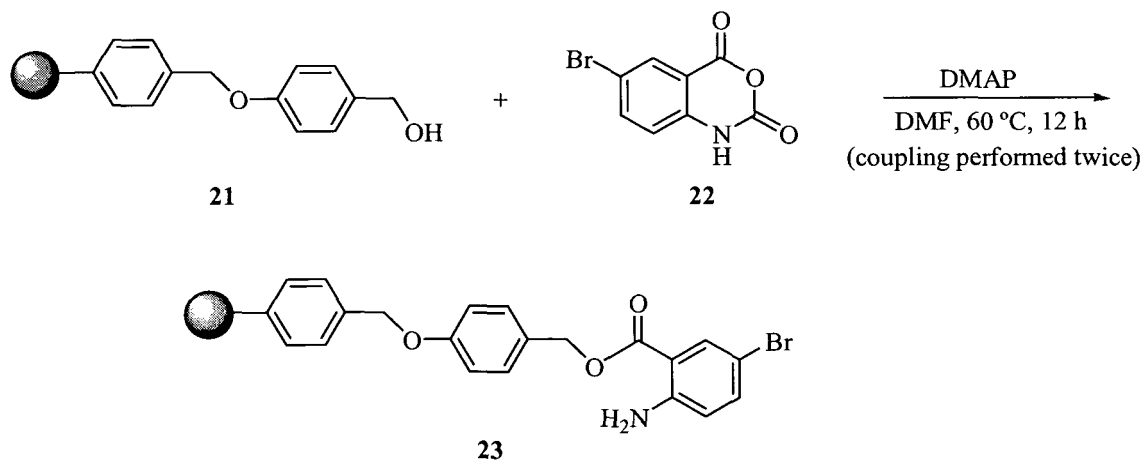
### Example 9: Library Synthesis

#### General Experimental

$^1\text{H}$  NMR spectra were measured using a Bruker AVANCE 300 spectrometer at rt in  $\text{DMSO}-d_6$  at an operating frequency of 300.13 MHz and are referenced to residual  $\text{DMSO}-d_6$  (2.54 ppm) unless otherwise noted. All coupling constants are reported in Hz. All non-combinatorial reactions were performed under a nitrogen atmosphere.

#### Synthetic Procedures Using Wang Resins

##### Scheme 9.1

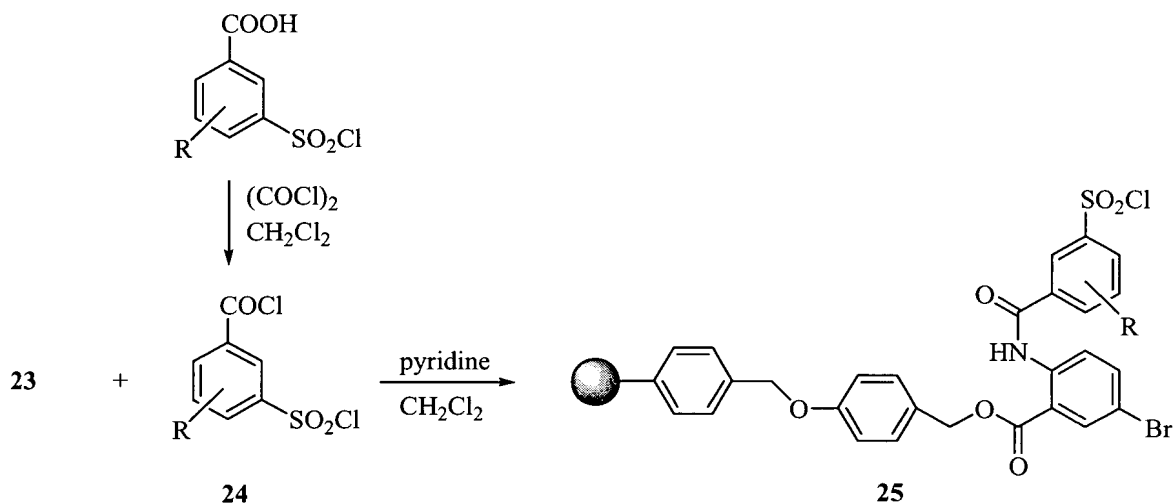


20

To a dry, 2-L polypropylene bottle equipped with a nitrogen inlet and an overhead stirrer was added Wang resin (21, 38.6 g, 49.7 mmol, 1.3 mmol/g, Novabiochem), DMF (600 mL), 5-bromoisatoic anhydride (22, 60.0 g, 248 mmol, dissolved in 100 mL of DMF), and DMAP (30.3 g, 248 mmol, dissolved in 100 mL of DMF). The reaction was heated under nitrogen to 65 °C and stirred for 12 h. The reaction was

then filtered and washed as follows: DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, and CH<sub>2</sub>Cl<sub>2</sub>. The washed resin was transferred back to the 2-L reaction flask and treated a second time with DMF (600 mL), 5-bromoisatoic anhydride (60.0 g, 248 mmol, dissolved in 100 mL of DMF), and DMAP (30.3 g, 248 mmol, dissolved in 100 mL of DMF). The reaction was stirred at 65 °C for 4 h and then filtered and washed with DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, and CH<sub>2</sub>Cl<sub>2</sub>) to afford (48.57 g) of **23** as an off-white resin. CNH analysis: Calcd (1.3 mmol): N, 1.82, Found: N, 1.67% (loading = 1.2 mmol/g).

10

**Scheme 9.2**

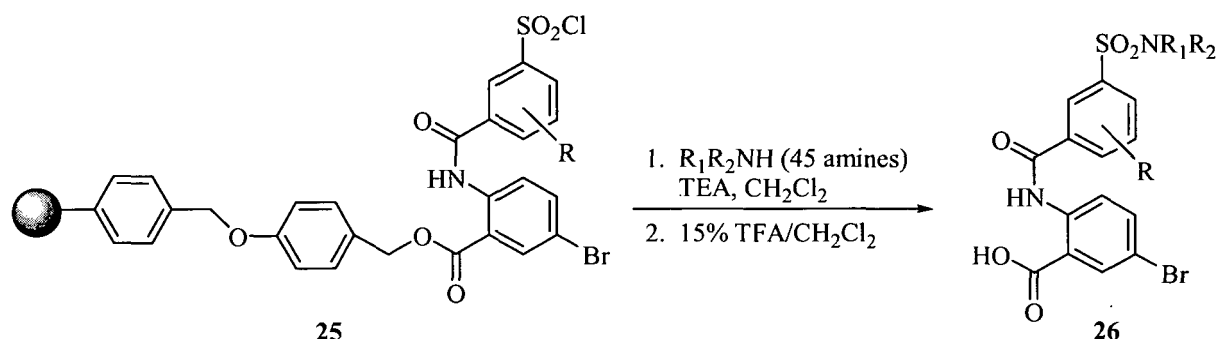
15

To a suspension of a 3-chlorosulfonylbenzoic derivative (**24**, 31.6 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (100 mL) was added DMF (two drops), followed by oxalyl chloride (31.6 mL of a 2 M solution in CH<sub>2</sub>Cl<sub>2</sub>, 63.2 mmol) under a nitrogen atmosphere. Gas evolution and disappearance of the suspension was noted during the course of the reaction. After the reaction was stirred for 18 h, the acid chloride was concentrated to dryness, azeotroped with toluene (2 x 25 mL), and then placed on a high vacuum. Dry anthranilic acid-derivatized Wang resin (7.0 g, 8.4 mmol) was added to an 8-oz wide-mouth bottle, followed by CH<sub>2</sub>Cl<sub>2</sub> (35 mL) and pyridine (35 mL). The acid chloride was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (20 mL) and added to resin, effecting HCl (g) evolution. The reaction jar was flushed with nitrogen, capped and shaken for 4 h.

25

The resin was then filtered and washed ( $\text{CH}_2\text{Cl}_2$ , MeCN,  $\text{CH}_2\text{Cl}_2$ , MeCN,  $\text{CH}_2\text{Cl}_2$ , MeCN,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CHCl}_3$ ,  $\text{CH}_2\text{Cl}_2$ , THF, MeCN, THF,  $\text{CH}_2\text{Cl}_2$ ; 50 mL each wash) to afford **25** as a tan resin.

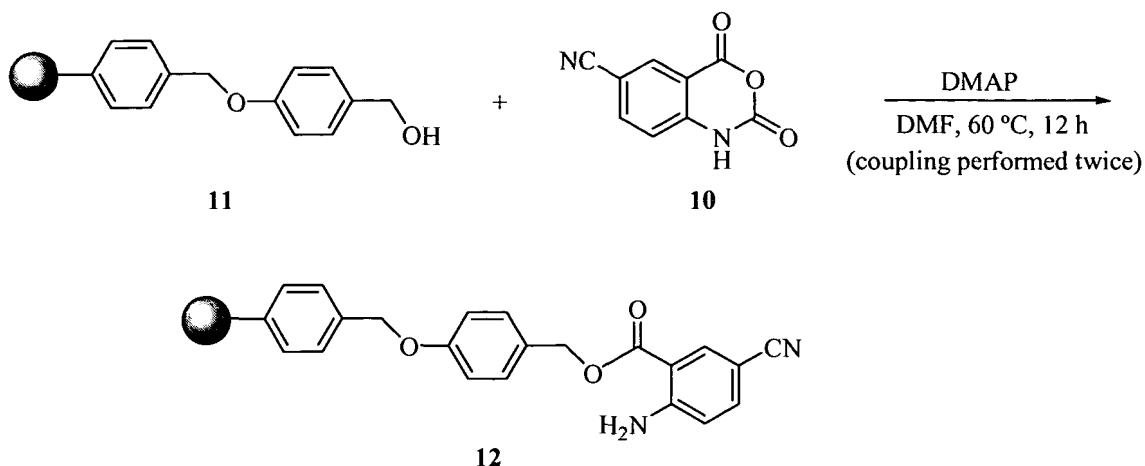
5 **Scheme 9.3**



Sulfonyl chloride resin (**25**, 50 mg, 60  $\mu\text{mol}$ ) was added down the columns of a 96-  
 10 well microtiter filter plate using a  $\text{CH}_2\text{Cl}_2$  isopycnic slurry. After draining the wells, the plate was inserted into a solid phase reaction apparatus. Amines (300  $\mu\text{L}$  of a 0.75 M solution, 225  $\mu\text{mol}$ ) were then added across the rows, followed by triethylamine (250  $\mu\text{L}$  of a 1.8 M solution) and  $\text{CH}_2\text{Cl}_2$  (250  $\mu\text{L}$ ). The plate was capped and spun on an overhead rotisserie for 16 h. After removal of the plate from the solid phase  
 15 reaction apparatus, the wells drained and each well was washed (DMF,  $\text{CH}_3\text{CN}$ , DMF,  $\text{CH}_3\text{CN}$ , DMF,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_3\text{CN}$ , and  $\text{CH}_2\text{Cl}_2$ ).

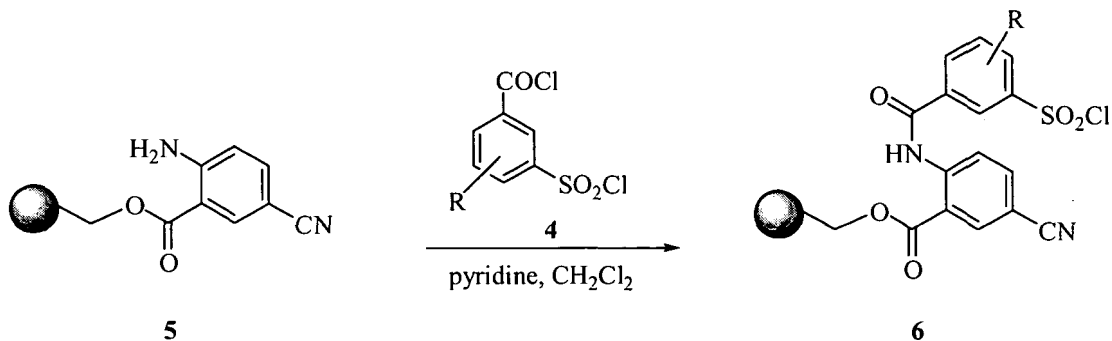
The plate was again inserted into the solid phase reaction apparatus and a 15%  
 20 solution of TFA in  $\text{CH}_2\text{Cl}_2$  (625  $\mu\text{L}$ ) was added. The plate was spun on an overhead rotisserie for 3 h and the crude sulfonamides were then drained into a 1-mL 96-well plate. The resin was washed with  $\text{CH}_2\text{Cl}_2$  (1.5 mL) and the washes collected in additional 1-mL plates. LC/MS samples were prepared by transferring 40  $\mu\text{L}$  of solution to a separate 96-well plate, concentrating the samples and then dissolving in  
 25 DMSO (125  $\mu\text{L}$ ) and diluting with acetonitrile (750  $\mu\text{L}$ ).

**Scheme 9.4**



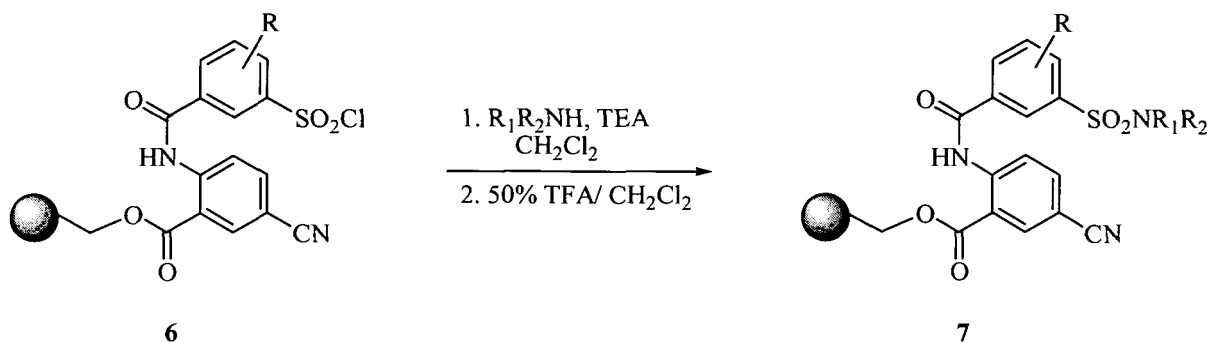
To a dry, 2-L polypropylene bottle equipped with a nitrogen inlet and an overhead stirrer was added Wang resin (**11**, 15.1 g, 21.1 mmol, 1.4 mmol/g, Novabiochem),  
 5  
 DMF (500 mL), 5-cyanoisatoic anhydride (**10**, 20.0 g, 106 mmol, dissolved in 100 mL DMF), and DMAP (13.0 g, 106 mmol, dissolved in 100 mL DMF). The mixture was heated under nitrogen to 53 °C and stirred for 16 h. The reaction was then filtered and washed with 500  $\mu$ L of the following solvents: DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF,  
 10  
 CH<sub>3</sub>CN, DMF, DMF, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, DMF, DMF, and DMF. The resin was transferred back to the 2-L reaction flask and treated a second time with DMF (500 mL), 5-cyanoisatoic anhydride (**10**, 20.0 g, 106 mmol, dissolved in 100 mL DMF), and DMAP (13.0 g, 106 mmol, dissolved in 100 mL DMF). The reaction was stirred at 60 °C for 22 h and then filtered and washed with 500  $\mu$ L of CH<sub>3</sub>CN, DMF,  
 15  
 CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, to afford 15.3 g of **12** as a pale yellow resin. Elemental analysis: N, 3.20 % (loading = 1.14 mmol/g).<sup>5</sup>

### Scheme 9.5



Dry 5-cyano anthranilic acid-derivatized Wang resin (**5**, 5.0 g, 1.0 mmol/g loading, 5.0 mmol) was added to an 8-oz wide mouth bottle, followed by  $\text{CH}_2\text{Cl}_2$  (30 mL) and pyridine (30 mL). The acid chloride (**4**) was dissolved in  $\text{CH}_2\text{Cl}_2$  (30 mL) and added to the resin, effecting HCl (gas) evolution. The jar was flushed with nitrogen, capped, and shaken for 64 h. The resin was then filtered and washed (DMF,  $\text{CH}_3\text{CN}$ , DMF,  $\text{CH}_3\text{CN}$ , DMF,  $\text{CH}_3\text{CN}$ , DMF, THF, THF, THF,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ; 400 mL each wash) to afford **6**.

### Scheme 9.6



Sulfonyl chloride resin (**6**, 50 mg, 50  $\mu\text{mol}$ ) was added to the wells of a 96-well filter plate using a  $\text{CH}_2\text{Cl}_2$  isopycnic slurry. After draining the wells, the plate was inserted into a solid phase reaction apparatus. Amines (250  $\mu\text{L}$  of a 2 M solution, 500  $\mu\text{mol}$ ) were then added, followed by triethylamine (250  $\mu\text{L}$  of a 2 M solution) and  $\text{CH}_2\text{Cl}_2$  (250  $\mu\text{L}$ ). The plate was then capped and spun on an overhead rotisserie for 20 h. After removal of the plate from the solid phase reaction block, the wells were drained and washed (DMF,  $\text{CH}_3\text{CN}$ , DMF,  $\text{CH}_3\text{CN}$ , DMF,  $\text{CH}_3\text{CN}$ ,  $\text{H}_2\text{O}$ , THF,  $\text{H}_2\text{O}$ , THF,  $\text{H}_2\text{O}$ , THF,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_3\text{CN}$ ,  $\text{CH}_2\text{Cl}_2$ ; 375  $\mu\text{L}$  each wash).

The plate was again inserted into the solid phase reaction apparatus and a 50% solution of TFA in  $\text{CH}_2\text{Cl}_2$  (500  $\mu\text{L}$ ) was added. The plate was spun on an overhead rotisserie for 3 h and the crude sulfonamides (**7**) were then drained into a standard 96-well plate. The resin was washed with 250  $\mu\text{L}$  of additional 50% TFA solution. Products were concentrated under nitrogen and then analyzed by LC/MS (see general LC/MS procedure).

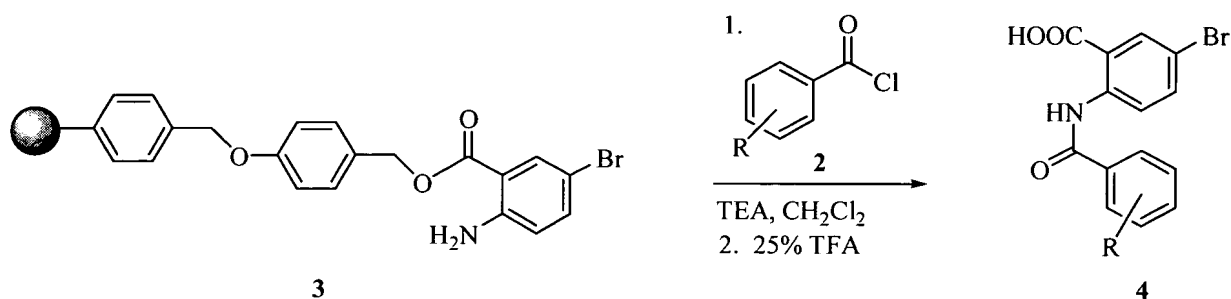


The crude samples were dissolved in THF, and eluted through a plug of Celite<sup>®</sup>.

LC/MS showed a reduced amount of impurity in all of the samples. The samples that were less than 70% pure were then eluted through a plug of silica gel using THF as the mobile phase and the samples were analyzed by LC/MS.

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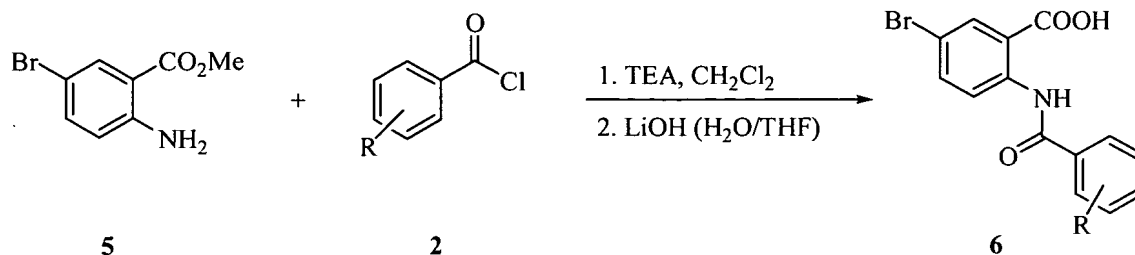
### Scheme 9.7



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To a standard 96-well filter plate was added 50 mg (60  $\mu$ mol) of 5-bromoanthranilic acid derivatized Wang resin as an isopycnic solution in CH<sub>2</sub>Cl<sub>2</sub> (3). After the wells were drained, the plate was inserted into a plate clamp assembly. The acid chloride diversity set (2) was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (300  $\mu$ L) and added to the plate, followed by TEA (250  $\mu$ L, 1 M CH<sub>2</sub>Cl<sub>2</sub>, 250  $\mu$ mol) and CH<sub>2</sub>Cl<sub>2</sub> (300  $\mu$ L). The plate was capped and spun on an overhead rotisserie for 16 h. After removal of the plate from the plate clamp assembly, the wells were drained and the resin washed with 500  $\mu$ L of the following solvents: CH<sub>2</sub>Cl<sub>2</sub>, MeCN, CH<sub>2</sub>Cl<sub>2</sub>, MeCN, CH<sub>2</sub>Cl<sub>2</sub>, MeCN, CH<sub>2</sub>Cl<sub>2</sub>, CHCl<sub>3</sub>, CH<sub>2</sub>Cl<sub>2</sub>, THF, MeCN, THF, CH<sub>2</sub>Cl<sub>2</sub>. The plate was reinserted into the plate clamp assembly and the washed resin was treated with 750  $\mu$ L of 25% TFA/CH<sub>2</sub>Cl<sub>2</sub> solution for 3 h. The solution was then filtered from the Wang resin and collected in a separate plate to afford the crude amides (4). The plates were concentrated and analyzed by LC/MS (see general LC/MS procedure).

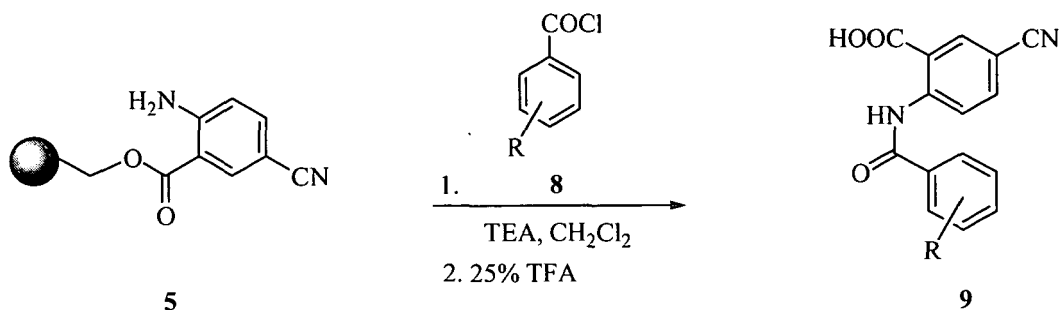
### 25 Scheme 9.8



After concentration of the acid chloride solutions (**2**), methyl-2-amino-5-bromobenzoate (**5**, 125  $\mu\text{L}$ , 1 M THF, 125  $\mu\text{mol/well}$ ) was added to the plate followed by potassium carbonate (1 mL, 0.38 M THF, 380  $\mu\text{mol/well}$ ). The reactions were capped, heated to 50  $^{\circ}\text{C}$  and shaken for 12 h. Triethylenetetramine resin (160 mg, 464  $\mu\text{mol}$ ) was added to the wells to scavenge the excess acid chloride and the plate spun for 2.5 h. The crude methyl esters were purified (if necessary) using a column consisting of basic alumina (ca 200 mg), SAX (ca 200 mg), and SCX (ca 400 mg, activated with 1% HOAc/MeOH) in descending order. The products were eluted with THF and the fractions analyzed by LC/MS.

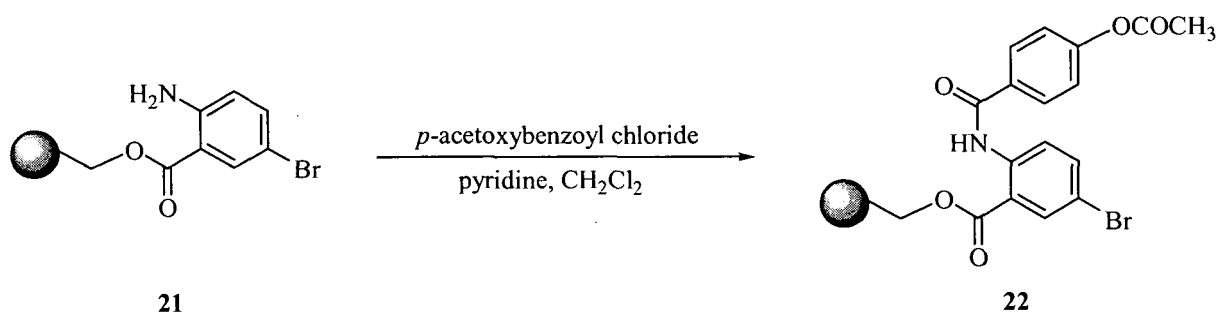
LiOH [375  $\mu\text{L}$ , 1 M  $\text{H}_2\text{O/THF}$  (50:50), 3 equiv)] was added to the esters and the plate was capped and spun for 1 h. The THF was then removed in vacuo. The crude solids were suspended in methyl ethyl ketone (MEK, 500  $\mu\text{L}$ ) and extracted with 2 N HCl (250  $\mu\text{L}$ ). The MEK layer was removed and the aqueous layer extracted again with MEK (500  $\mu\text{L}$ ). The combined organic layers were washed with 50% brine solution, passed through a plug of sodium sulfate, collected in a 1-mL plate, and dried under nitrogen to afford the amide products (**6**). The solids were then analyzed using LC/MS (see general LC/MS procedure).

### Scheme 9.9



To each vial of an array of 1-mL vials arranged in a 96-well format was added 44 mg (50  $\mu$ mol) of 5-cyanoanthranilic acid-derivatized Wang resin (**5**) as an isopycnic solution in CH<sub>2</sub>Cl<sub>2</sub>. The acid chloride diversity set<sup>2</sup> (**8**, 500  $\mu$ mol) was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (300  $\mu$ L) and added to the vials, followed by TEA (250  $\mu$ L, 2 M CH<sub>2</sub>Cl<sub>2</sub>, 500  $\mu$ mol), and CH<sub>2</sub>Cl<sub>2</sub> (300  $\mu$ L). The vials were capped, heated to 60 °C, and shaken for 21 h. After completion of the reaction, the resin was transferred to a 96-well filter plate and washed with of the following solvents: DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, H<sub>2</sub>O, THF, H<sub>2</sub>O, THF, H<sub>2</sub>O, THF, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub> (500  $\mu$ L/wash). The plate was placed into a clamp assembly and each well was treated with 500  $\mu$ L of 50% TFA/CH<sub>2</sub>Cl<sub>2</sub> solution for 2 h.<sup>3</sup> The resultant solution was then filtered from the Wang resin, collected in a separate plate, and dried under nitrogen to afford the crude amides (**9**).

#### Scheme 9.10

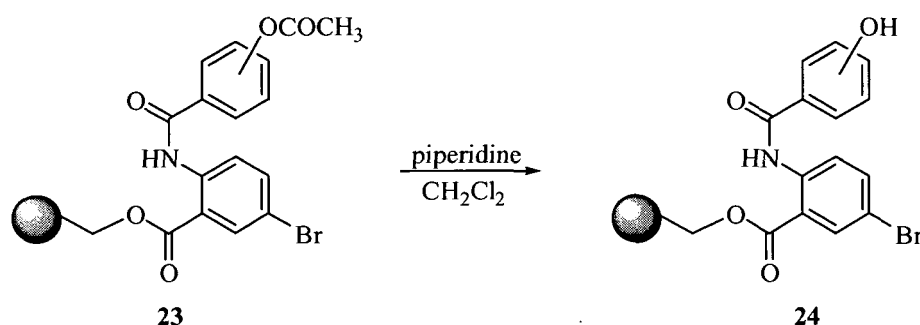


**Resin-bound 4-Acetoxybenzoyl Anthranilic Acid.** To a 500-mL round bottom flask under nitrogen was added 4-acetoxybenzoic acid (20.7 g, 115.5 mmol) and CH<sub>2</sub>Cl<sub>2</sub> (200 mL). After cooling the flask to 0 °C, oxalyl chloride (57.8 mL of a 2 M solution, 116 mmol) and a few drops of DMF were added. The reactions were allowed to warm to room temperature and stirred for 3 h. These solutions were directly transferred to a

2-L serum flask containing 5-bromoanthranillic acid resin (**21**, 7.0 g, 7.7 mmol), pyridine (100 mL) and CH<sub>2</sub>Cl<sub>2</sub> (100 mL). The resulting mixtures were stirred under nitrogen overnight and then filtered into a glass fritted funnel. The resin was then washed with DMF (3 x 100 mL), CH<sub>2</sub>Cl<sub>2</sub> (5 x 100 mL), and MeOH (5 x 100 mL).

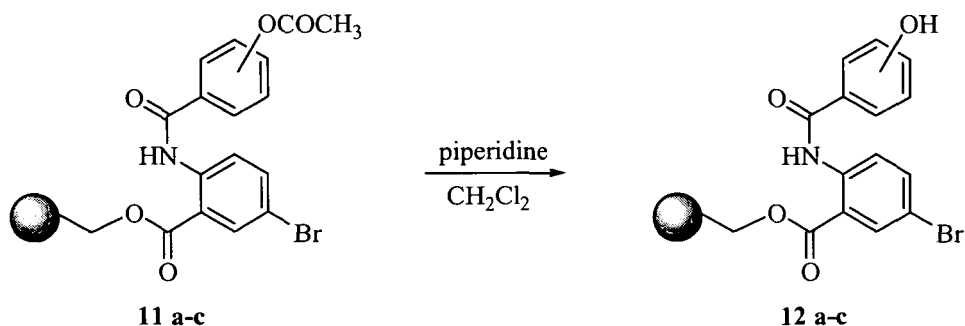
- 5 The resin was then dried in a vacuum oven at 60 °C for 72 h to afford **22** (8.0 g). A sample was cleaved from the resin by stirring in 25% TFA in CH<sub>2</sub>Cl<sub>2</sub> for 3 h: <sup>1</sup>H NMR (acetone-*d*<sub>6</sub>) δ 2.31 (s, 3H), 7.35 (d, *J* = 2.1, 1H), 7.37 (d, *J* = 2.0, 1H), 7.82 (d, *J* = 2.5, 1H), 7.86 (d, *J* = 2.5, 1H), 8.07 (dd, *J* = 2.1, 8.7, 1H), 8.27 (d, *J* = 2.5, 1H), 8.90 (d, *J* = 9.0, 1H).

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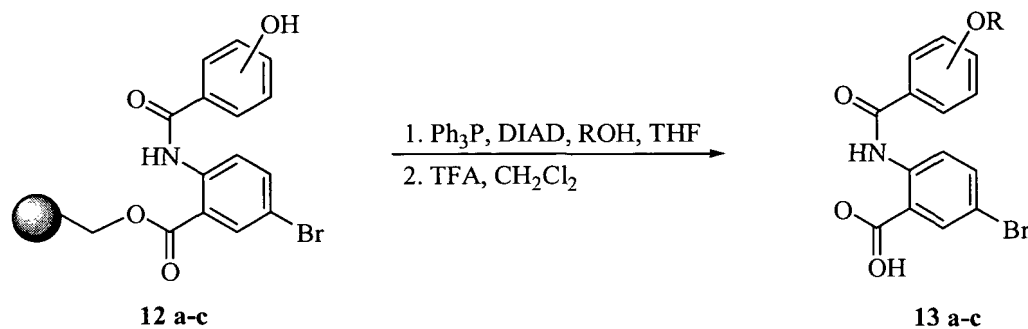
- Resin-bound 4-Hydroxybenzoyl Anthranilic Acid.** To a 250-mL serum bottle was added acetoxy resin **23** (7.0 g, 7.7 mmol), CH<sub>2</sub>Cl<sub>2</sub> (70 mL), and piperidine (150 mL, 2 M CH<sub>2</sub>Cl<sub>2</sub>). The slurry was stirred for 2 h at room temperature. The resins were then filtered and washed with DMF (3 x 100 mL), Et<sub>3</sub>N (1 M CH<sub>2</sub>Cl<sub>2</sub>, 2 x 100 mL), and MeOH (2 x 100 mL), CH<sub>2</sub>Cl<sub>2</sub> (40 mL), MeOH (40 mL), CH<sub>2</sub>Cl<sub>2</sub> (40 mL), MeOH (40 mL), CH<sub>2</sub>Cl<sub>2</sub> (40 mL), and MeOH (40 mL). The resin was then dried for 72 h in a vacuum oven at room temperature to afford 6.6 g of **24** as a yellow resin.

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**Synthesis of Resin-Bound Phenol 12a.** To a 200-mL Wheaton bottle equipped with an overhead stirrer was added resin-bound acetate (**11a**, 5.0 g) followed by piperidine (150 mL of a 2 M solution in CH<sub>2</sub>Cl<sub>2</sub>, 300 mmol). The reaction was stirred for 2 h at room temperature. The resin was then filtered from the reaction mixture, washed with

5 DMF, DMF, DMF, Et<sub>3</sub>N (1 M in CH<sub>2</sub>Cl<sub>2</sub>), MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, (50 mL each), and dried in a vacuum oven at 50 °C overnight to afford resin-bound phenol **12a** as a brownish solid.



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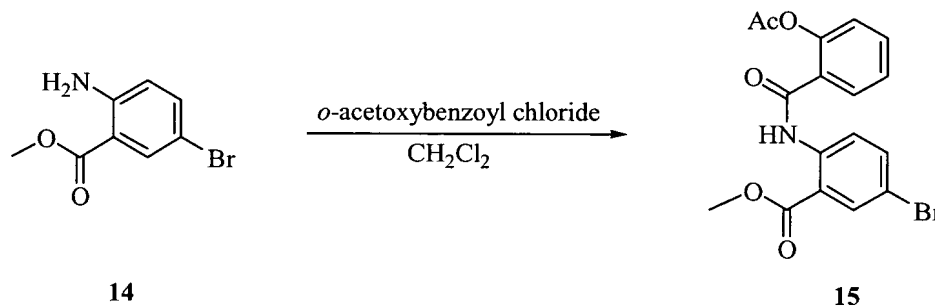
**Mitsunobu Reaction (Procedure A).** To each well of a fritted 96-well plate was added phenol resin (**12a-c**, 20.0 mg, 20.0 μmol) as an isopycnic solution (20% THF in CH<sub>2</sub>Cl<sub>2</sub>) and the plate was placed in a solid phase reaction assembly. The alcohol diversity element (200 μL of a 1 M solution in THF, 200 μmol) was then added,

15 followed by triphenylphosphine (200 μL of a 1 M solution in THF, 200 μmol). The wells were flushed with nitrogen, capped, and placed in the -20 °C freezer for 1 h. While in the freezer, DIAD [200 μL of a cooled (-20 °C), freshly made 1 M solution in THF] was added to each well. The plate was removed from the freezer after 1 h and then spun on the rotisserie for 16 h. The reaction mixture was drained from the

20 plate and the resin then washed with THF, THF, THF (the plate was capped and spun on an overhead rotisserie for 30 min), THF, MeOH, THF, MeOH, THF, MeOH, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH, CH<sub>2</sub>Cl<sub>2</sub>, MeOH (the plate was capped and spun on an overhead rotisserie for 30 min), CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>; 500 μL each solvent. The crude aryl ethers were

25 then cleaved from the resin using 500 μL of 50% TFA in CH<sub>2</sub>Cl<sub>2</sub>. The resulting products (**13a-c**) were concentrated under a nitrogen stream and analyzed by HPLC/MS.

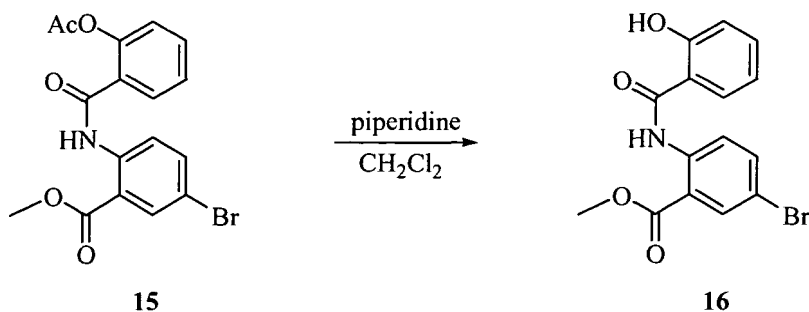
**Mitsunobu Reaction (Procedure B).** To 72 wells of a fritted 96-well plate was added phenol resin as an isopycnic solution (**12a-c**, 20.0 mg, 20.0  $\mu\text{mol}$ ) and the plate was placed in a solid phase reaction assembly. The alcohol diversity element (200  $\mu\text{L}$  of a 1 M solution in THF, 200  $\mu\text{mol}$ ) was then added, followed by triphenylphosphine (200  $\mu\text{L}$  of a 1 M solution in THF, 200  $\mu\text{mol}$ ) and  $\text{Et}_3\text{N}$  (200  $\mu\text{L}$  of a 1 M solution in THF, 200  $\mu\text{mol}$ ). The wells were flushed with  $\text{N}_2$ , capped, and placed in the  $-20^\circ\text{C}$  freezer for 1 h. While in the freezer, DIAD [200  $\mu\text{L}$  of a cooled ( $-20^\circ\text{C}$ ), freshly made 1 M solution in THF] was added to each well. The plate was removed from the freezer after an hour and then spun on the rotisserie for 16 h. The reaction mixture was drained from the plate and the resin then washed with THF, THF, THF (the plate was capped and spun on an overhead rotisserie for 30 min), THF, MeOH, THF, MeOH, THF, MeOH, MeOH,  $\text{CH}_2\text{Cl}_2$ , MeOH,  $\text{CH}_2\text{Cl}_2$ , MeOH,  $\text{CH}_2\text{Cl}_2$ , MeOH,  $\text{CH}_2\text{Cl}_2$ , MeOH,  $\text{CH}_2\text{Cl}_2$ , MeOH,  $\text{CH}_2\text{Cl}_2$ , MeOH (the plate was capped and spun on an overhead rotisserie for 30 min),  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CH}_2\text{Cl}_2$ ; 500  $\mu\text{L}$  each solvent. The crude aryl ethers were then cleaved from the resin using 500  $\mu\text{L}$  of 50% TFA in  $\text{CH}_2\text{Cl}_2$ . The resulting products (**13a-c**) were concentrated under a nitrogen stream and analyzed by HPLC/MS.



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**Synthesis of Acetate 15.** To a 250-mL round bottom flask was added a solution of methyl-2-amino-5-bromobenzoate [**14**, 5.0 g, 21.7 mmol dissolved in pyridine (10 mL) and  $\text{CH}_2\text{Cl}_2$  (10 mL)], followed by *o*-acetoxybenzoyl chloride<sup>5</sup> (4.7 g, 33.8 mmol dissolved in 60 mL of  $\text{CH}_2\text{Cl}_2$ ). The mixture was stirred overnight under a nitrogen atmosphere. Polyamine resin (4.0 g) was then added to the reaction mixture and the reaction was stirred for 4 h. After filtration and concentration of the reaction mixture, a white residue was obtained. The residue was recrystallized from  $\text{CH}_2\text{Cl}_2$  to afford 8.0 g (94%) of **15** as a white solid:  $^1\text{H}$  NMR ( $\text{DMSO}-d_6$ )  $\delta$  2.24 (s, 3H), 3.86 (s, 3H),

7.30 (d,  $J = 8.1$ , 1H), 7.46 (dt,  $J = 1.1$ , 7.6, 1H), 7.66 (dt,  $J = 1.7$ , 8.0, 1H), 7.82 (dd,  $J = 1.7$ , 7.7, 1H), 7.86 (dd,  $J = 2.5$ , 8.9, 1H), 8.06 (d,  $J = 2.5$ , 1H), 8.38 (d,  $J = 8.9$ , 1H).

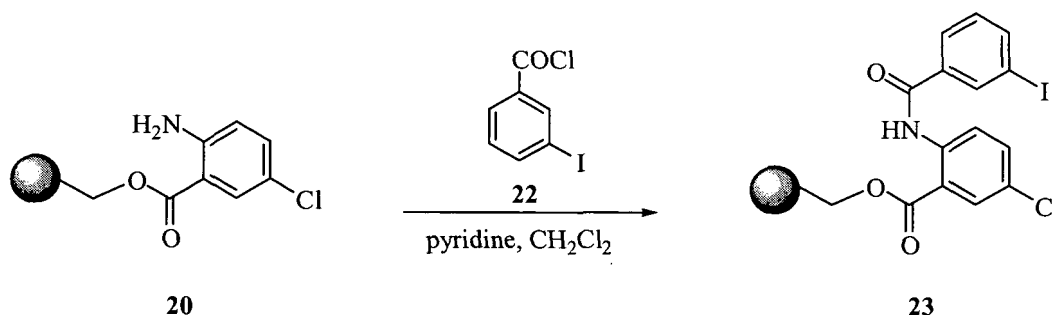


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**Synthesis of Phenol 16.** To a 50-mL round bottom flask was added *o*-acetoxy methyl ester **15** (1.0 g, 2.9 mmol), CH<sub>2</sub>Cl<sub>2</sub> (10 mL) and piperidine (2.0 mL of a 2 M solution in CH<sub>2</sub>Cl<sub>2</sub>, 4.0 mmol). After the reaction mixture was stirred for 3 h, the solvent was removed and the crude residue dried under high vacuum overnight. The residue was then dissolved in CH<sub>2</sub>Cl<sub>2</sub> and acid chloride resin (2.0 g, 2.1 mmol) was added to scavenge excess piperidine. The mixture was stirred for 4 h, filtered, and concentrated to afford 0.54 g (60%) of phenol **16** as a white solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>)  $\delta$  3.90 (s, 3H), 6.98 (t,  $J = 7.6$ , 1H), 7.02 (d,  $J = 7.6$ , 1H), 7.44 (dt,  $J = 1.8$ , 8.2, 1H), 7.82 (dd,  $J = 2.5$ , 9.0, 1H), 7.93 (dd,  $J = 1.8$ , 7.9, 1H), 8.08 (d,  $J = 2.5$ , 1H), 8.60 (d,  $J = 9.0$ , 1H).

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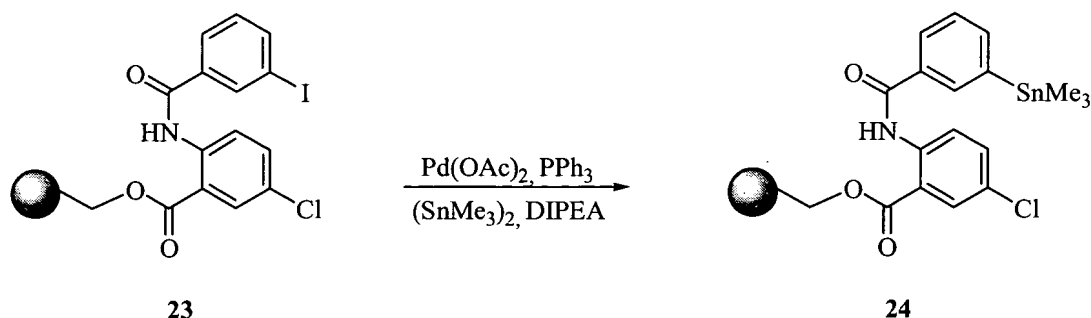
### Scheme 9.11



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**Resin-Bound *m*-Iodo Benzamide 23.** Acid chloride **22** was redissolved in CH<sub>2</sub>Cl<sub>2</sub> (30 mL) and added to resin-bound 5-chloroanthranilic acid (**20**, 3 g, 1.06 mmol/g loading, 3.18 mmol) swollen with pyridine (30 mL) in a 500-mL serum flask equipped with an overhead stirrer. The flask was purged with nitrogen and the resin stirred for

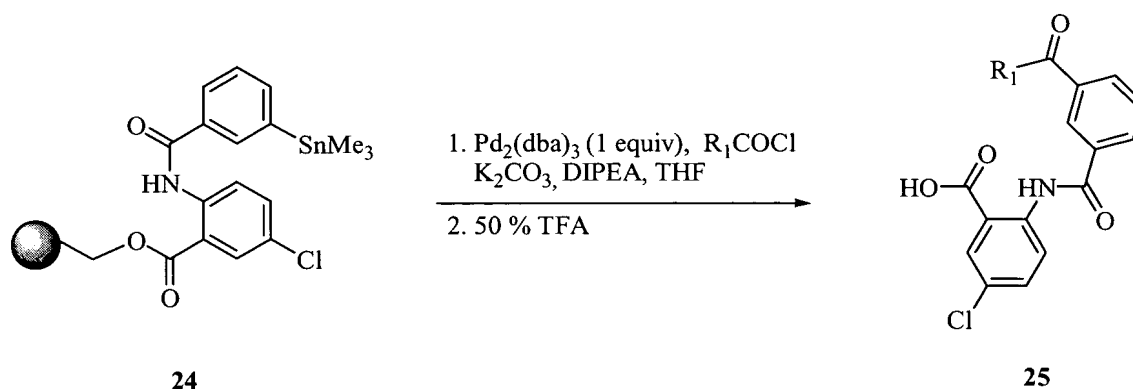
16 h. The resin was filtered from the reaction mixture and washed with alternating  $\text{CH}_3\text{CN}$  and  $\text{CH}_2\text{Cl}_2$  washes (8 x 300 mL) to afford **23**.



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**Resin-Bound Stannylate 24.** To a  $\text{CH}_2\text{Cl}_2$  slurry of *m*-iodo resin (**23**, 1 g, 1.06 mmol/g) in a 250-mL serum flask was added 1 mL of the following solutions; palladium acetate (0.0022 g/ 1 mL, 0.01 mmol, 0.1 equiv.), triphenyl phosphine (0.0065 g/mL, 0.025 mmol, 0.25 equiv), DIPEA (0.0065 g/mL, 0.05 mmol, 0.5 equiv) in DMF. Hexamethyl ditin (0.065 g, 0.2 mmol, 2.0 equiv) was added to the flask, which was then purged with nitrogen and heated to 60 °C for 18 h. The reaction mixture was drained and the resin washed with alternating DMF,  $\text{CH}_3\text{CN}$  and  $\text{CH}_2\text{Cl}_2$  (10 x 150 mL) to yield **24** as a dark brown resin.

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**Resin Bound Library of Aryl Ketones.** Hexamethyl ditin derivatized Wang resin (**24**, 24 mg, 24  $\mu\text{mol}$ ) was added as an isopycnic solution (degassed THF) to an array of 1-dram vials arranged in a 96-well format. Tris(dibenzylidene acetone) dipalladium (0) (22 mg, 24  $\mu\text{mol}$ , 1.0 equiv) was added to each vial (in a solution of degassed THF). DIPEA (20  $\mu\text{L}$ ) was added to each vial followed by  $\text{K}_2\text{CO}_3$  (10 mg)

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and degassed THF (0.5 mL). The vials were capped and shaken. The vials were uncapped and the acid chloride diversity elements<sup>7</sup> (10 equiv) were then added, the vials purged with nitrogen for 5 sec, capped, shaken and heated 60 °C for 20 h. After the reactions cooled to room temperature, the resin was transferred to a 96-well polypropylene fritted plate. The resin was washed (CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF, CH<sub>3</sub>CN, DMF, H<sub>2</sub>O, THF, H<sub>2</sub>O, THF, H<sub>2</sub>O, THF, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>3</sub>CN, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, 250 µL each wash) and the plate inserted into a solid phase reaction block. A solution of 50% TFA in CH<sub>2</sub>Cl<sub>2</sub> (600 µL) was added to the plate. The plate was capped and spun on an overhead rotisserie for 3 h. The crude aryl ketones (**25**) were then drained into a 96-well collection plate, concentrated to dryness, and analyzed by HPLC/MS.

### Purification Procedures

**Liquid-liquid extraction (basic).** To a 96-well plate of crude samples was added methyl ethyl ketone (MEK, 500 µL) and 2 N NaOH (500 µL). The plates were capped and shaken. After the plates were uncapped, the organic layer was separated from the aqueous layer.

**Liquid-liquid extraction (acidic).** The aqueous layer of the above extraction was treated with 6 N HCl (500 µL) and extracted with MEK (1 mL). The plates were capped, shaken, and the organic layer was separated from the aqueous layer.

**Hydromatrix<sup>®</sup> extraction** (AMRI SEC-C-44). A set of 2-mL square-well plates were filled with Hydromatrix<sup>®</sup> and washed with MEK and CH<sub>2</sub>Cl<sub>2</sub> (500 µL/well). The plates were then placed in a vacuum oven (T = 35 °C) overnight. After cooling, the Hydromatrix<sup>®</sup> was treated with 2 N HCl (600 µL)<sup>7</sup> and the plates were stacked. The crude library samples were dissolved in MEK and pipetted onto the columns. MEK was used to elute the compounds, and several 2-mL fractions were collected.

### Crystallization

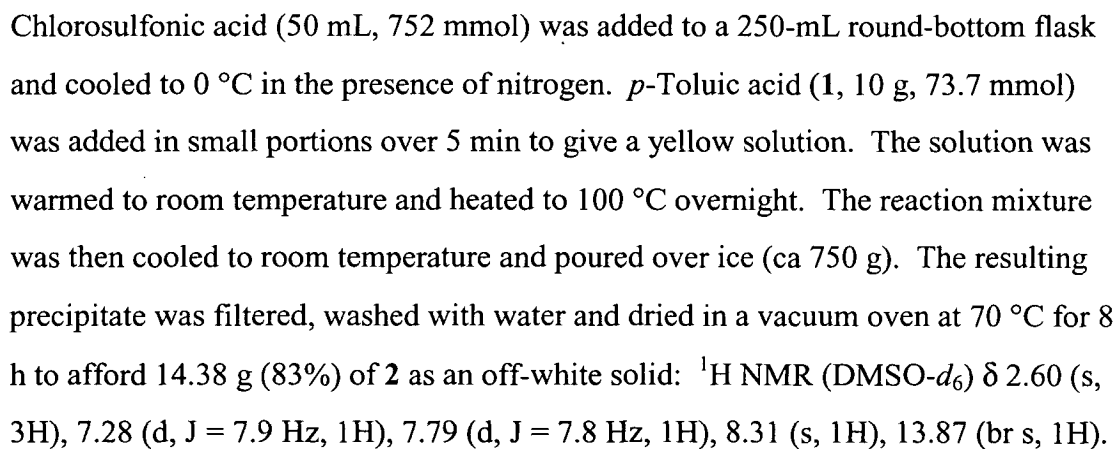
After treatment with Hydromatrix<sup>®</sup>, several compounds crystallized out of the 50% MeOH/MEK solution. The liquid was removed from the well, and the solid dissolved in DMSO (250 µL) and transferred to a Marsh tube.

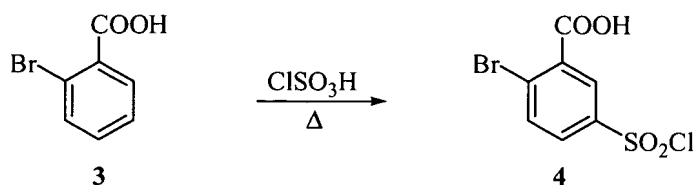
- 5    **HPLC analysis method.** The purity of the library was determined from the relative peak area of the UV absorbance. The identity of the compound was determined by MS confirmation of the molecular weight. The samples from this library were best prepared from DMSO solutions of the crude compounds. To a 96-well LC/MS plate was added ca 30 µL of DMSO solution (solution concentration was typically ca 30
- 10    mM). DMSO (ca 50 µL) and MeCN (ca 750 µL) were then used to dilute the samples.

#### HPLC Conditions

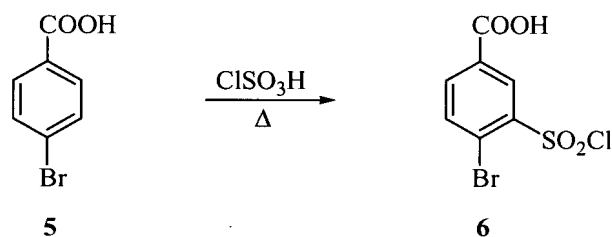
- 15    Column:                    Zorbax SB-C18 (4.6 x 75 mm, 3.5 microns)  
Gradient: A solvent: 100% MeCN (0.075% HCO<sub>2</sub>H), B solvent: 100% H<sub>2</sub>O (0.075% HCO<sub>2</sub>H)  
Flow:                        2 mL/min  
Detection wavelength: 220 nm (UV)
- 20    Autosampler:              Gilson 215 Liquid Handler  
Pump:                        Shimadzu LC-10AD VP  
Detector:                    Shimadzu UV-VIS Detector SPD-10A VP  
Injection volume:        40 µL  
Mass Spectrometer:      PESCIEX API 150EX

### Preparation of Benzoic Acid Derivatives for Library Synthesis

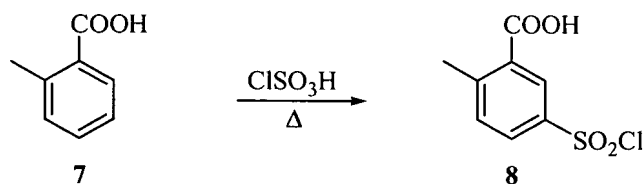
10



To a 250-mL round-bottom flask cooled to 0 °C under nitrogen was added  
 5 chlorosulfonic acid (50 mL, 752 mmol), followed by *o*-bromobenzoic acid (**3**, 10.0 g, 49.7 mmol) in small portions over 2 min to give a brownish solution. The solution was warmed to room temperature and heated to 115 °C overnight. The reaction mixture was then cooled to room temperature and poured over ice (ca 750 g).<sup>1</sup> The resulting precipitate was filtered, washed with water and dried in a vacuum oven at 80  
 10 °C for 7 h to afford 12.81 g (86%) of **4** as an off-white solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 7.75 (d, *J* = 10.1 Hz, 1H), 7.65 (d, *J* = 10.1 Hz, 1H), 8.46 (s, 1H), 13.96 (br s, 1H).

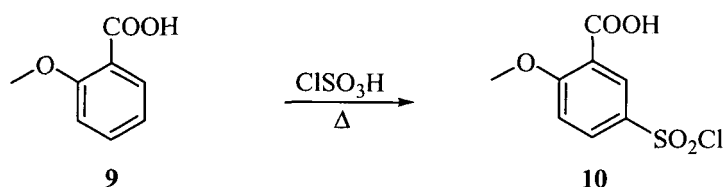


15 Chlorosulfonic acid (50 mL 752 mmol) was added to a 250-mL round-bottom flask and cooled to 0 °C in the presence of nitrogen. *p*-Bromobenzoic acid (**5**, 10.0 g, 49.7 mmol) was added in small portions over 2 min to give a brownish solution. The solution was warmed to room temperature and heated to 145 °C overnight. The reaction mixture was then cooled to room temperature and poured over ice (ca 750  
 20 g).<sup>1</sup> The resulting precipitate was filtered, washed with water and dried in a vacuum oven at 80 °C for 7 h to afford 13.21 g (89%) of **6** as a tan solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 7.60 (dd, *J* = 2.1, 8.3 Hz, 1H), 7.69 (d, *J* = 8.2 Hz, 1H), 8.31 (d, *J* = 2.1 Hz, 1H), 14.05 (br s, 1H).

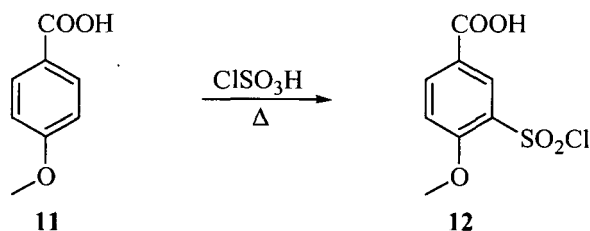


Chlorosulfonic acid (50 mL, 752 mmol) was added to a 250-mL round-bottom flask and cooled to 0 °C in the presence of nitrogen. *o*-Toluic acid (**7**, 10.0 g, 73.4 mmol) was added in small portions over 2 min to give a brownish solution. The solution was  
 5 warmed to room temperature and heated to 145 °C overnight. The reaction mixture was then cooled to room temperature and poured over ice (ca 750 g).<sup>1</sup> The resulting precipitate was filtered, washed with water and dried in a vacuum oven at 80 °C for 7 h to afford 15.53 g (90%) of **8** as an off-white solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 2.53 (s, 3H), 7.26 (d, *J* = 7.9 Hz, 1H), 7.63 (d, *J* = 7.9 Hz, 1H), 8.07 (s, 1H), 13.60 (br s, 1H).

10



Chlorosulfonic acid (50 mL, 752 mmol) was added to a 250-mL round-bottom flask and cooled to 0 °C in the presence of nitrogen. *p*-Anisic acid (**9**, 10.0 g, 73.4 mmol) was added in small portions over 2 min to give a yellow solution. The solution was  
 15 warmed to room temperature and heated to 63 °C for 1 h. The reaction mixture was then cooled to room temperature and poured over ice (ca 750 g).<sup>1</sup> The resulting precipitate was filtered, washed with water and dried in a vacuum oven at 70 °C for 12 h to afford 14.62 g (85%) of **10** as a white solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 3.84 (s, 3H), 7.06 (d, *J* = 8.7 Hz, 1H), 7.70 (dd, *J* = 2.3, 8.7 Hz, 1H), 8.31 (d, *J* = 2.3 Hz, 1H),  
 20 13.82 (br s, 1H).

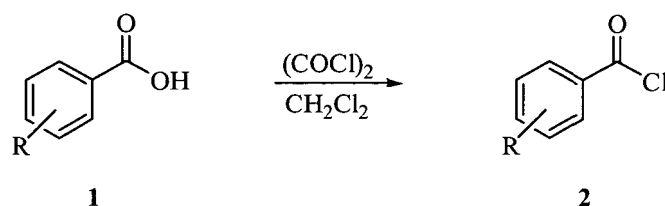


Solid *p*-anisic acid (**11**, 10.0 g, 66 mmol) was added to an ice-cooled, 250-mL round-bottom flask containing chlorosulfonic acid (50 mL, 752 mmol) under nitrogen. The  
 25 solution was heated at 65 °C for 1 h and turned bright yellow. The reaction mixture was cooled to room temperature and poured over ice (ca 750 g). The resulting

precipitate was then filtered, washed with water and dried in a vacuum oven at 70 °C for 8 h to yield 13.18 g (80%) of **12** as a pale yellow solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 3.88 (s, 3H), 7.06 (d, *J* = 8.7 Hz, 1H), 7.90 (dd, *J* = 2.4, 8.6 Hz, 1H), 8.31 (d, *J* = 2.3 Hz, 1H), 13.82 (br s, 1H).

5

### General Procedure for the Conversion of Acids to Acid Chlorides in a Plate Format

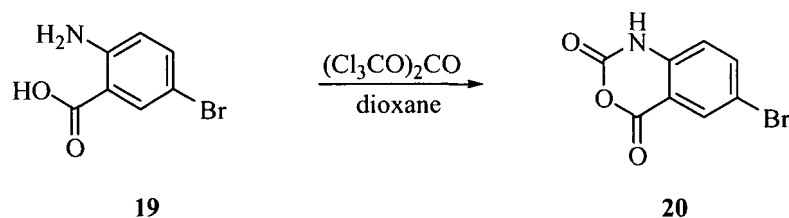


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To a plate of 2-mL glass reaction tubes arranged in a standard 96-well format was added the diversity set of carboxylic acids (**1**, 250 μL, 1 M THF, 250 μmol). The samples were concentrated in a Genevac HT-4 (20% heat with no heat boost for 1 h). A solution of 1% DMF/CH<sub>2</sub>Cl<sub>2</sub> (50 μL) was added to the wells, followed by CH<sub>2</sub>Cl<sub>2</sub> (250 μL). The carboxylic acid plate was placed in a nitrogen-filled glove bag and oxalyl chloride (125 μL, 2 M CH<sub>2</sub>Cl<sub>2</sub>, 250 μmol) was added. After the addition of CH<sub>2</sub>Cl<sub>2</sub> (250 μL), a capmat with 96 predrilled holes was fitted on the plate. The plate was shaken on an orbital shaker in a N<sub>2</sub> filled glove bag for 6-8 h.

15

### 20 Preparation of Isatioc Anhydride Derivatives



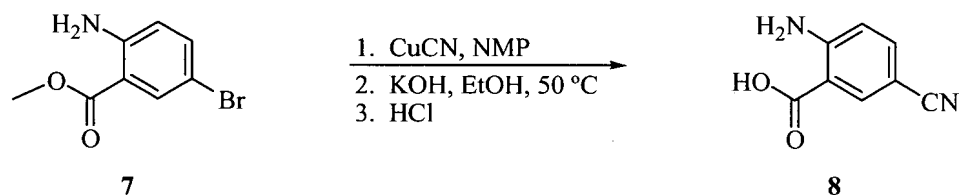
To a dry, 4-L round bottom flask was added 175 g (810 mmol) of 2-amino-5-bromobenzoic acid (**19**), triphosgene (83 g, 278 mmol), and dioxane (3 L). The suspension was stirred under N<sub>2</sub> and heated to reflux. The reaction was found to be complete by TLC and NMR after stirring at reflux for 3 h, but did not become homogenous at any time. After cooling to room temperature, the reaction was filtered

25

and the precipitate washed with ether. The solid was dried in the vacuum oven at 40 °C to afford 5-bromoisatoic anhydride (**20**, 151.1 g, 72%) as a white solid:  $^1\text{H}$  NMR (DMSO- $d_6$ )  $\delta$  7.29 (d,  $J$  = 8.7, 1H), 7.91 (dd,  $J$  = 2.5, 8.7, 1H), 8.09 (d,  $J$  = 2.3, 1H).

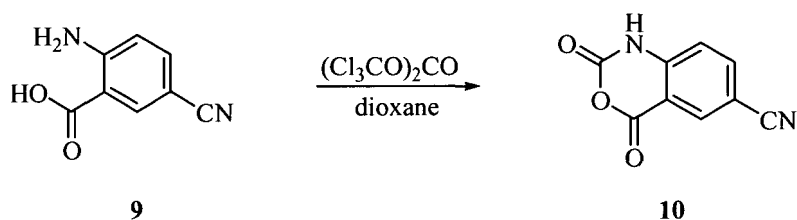
- 5 Sodium cyanoborohydride (4.88 g, 77.8 mmol) was added to a solution of 6-chloroindoline (5.9 g, 38.9 mmol) in acetic acid (100 mL). Gas evolution was evident at the beginning of the reaction. After stirring for 10 h, the solution was diluted with water (100 mL) and 6 N NaOH was added until the pH of the reaction mixture was 12-13. The resulting mixture was extracted with  $\text{CH}_2\text{Cl}_2$  (3 x 200 mL), and the
- 10 combined organic layers dried over  $\text{MgSO}_4$ . Flash column chromatography on silica gel (35% EtOAc/hexanes) yielded 2.3 g (39%) of a clear liquid:  $^1\text{H}$  NMR (DMSO- $d_6$ )  $\delta$  2.87 (t,  $J$  = 8.4 Hz, 2H), 3.44 (t,  $J$  = 8.4 Hz, 2H), 6.45 (d,  $J$  = 1.8 Hz, 1H), 6.47 (dd,  $J$  = 1.8, 7.6 Hz, 1H), 6.96 (d,  $J$  = 7.3 Hz, 1H).

15

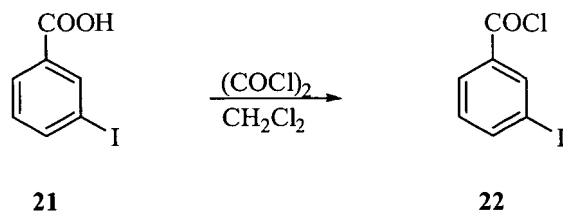


- To a 3-L, three-necked, round bottom flask equipped with a reflux condenser was added methyl-2-amino-5-bromobenzoate (**7**, 125 g, 543 mmol), copper cyanide (56.2 g, 624 mmol), and NMP (1 L). The reaction was heated to 200 °C and stirred for 4 h
- 20 under nitrogen. The dark brown reaction mixture was allowed to cool and a brown precipitate was formed. The mixture was poured into a 16-L beaker containing sodium cyanide solution (1 kg NaCN in 6 L  $\text{H}_2\text{O}$ ) followed by the addition of EtOAc (4 L). The precipitate was dissolved by agitation and the layers were separated. The
- 25 aqueous layer was extracted with EtOAc (2 x 1.5 L) and the combined organic layers were washed with 10% NaCN solution (2 L),  $\text{H}_2\text{O}$  (2 L) and then dried over  $\text{MgSO}_4$ . The light brown solution was concentrated and then dried in a vacuum oven overnight.

The ester was then dissolved in EtOH (2 L) and added to a 3-L round bottom flask followed by KOH solution (96.7 g of KOH in 500 mL H<sub>2</sub>O). The reaction mixture was heated to 50 °C and stirred for 2 h. The resultant dark-brown solution was poured into a chilled 2 N HCl solution (1.5 L), creating a yellowish precipitate. The solid was collected on a sintered glass filter frit, washed with cold water, and dried at 35 °C in a vacuum oven overnight to give 64.0 g (73%) of 5-cyanoanthranilic acid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 7.16 (d, *J* = 8.7, 1H), 7.79 (dd, *J* = 2.5, 8.7, 1H), 7.87 (d, *J* = 2.4, 1H).



To a dry 4-L round bottom flask was added 5-cyanoanthranilic acid (**9**, 64 g, 395 mmol), triphosgene (39.4 g, 131 mmol) and dioxane (2 L). The suspension was stirred under N<sub>2</sub> and heated to reflux. The reaction mixture became homogeneous after stirring at reflux for 2 h. As the carbonylation product was formed, white precipitate appeared in the solution. After stirring at reflux for an additional 3 h, the reaction was cooled to room temperature, filtered, and the precipitate washed with ether. The solid was dried in the vacuum oven to afford 5-cyanoisatoic anhydride (**10**, 51.5 g, 68%) as a pale yellow solid: <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>) δ 6.86 (d, *J* = 9.3, 1H), 7.55 (dd, *J* = 2.6, 9.0, 1H), 8.04 (d, *J* = 2.4, 1H).



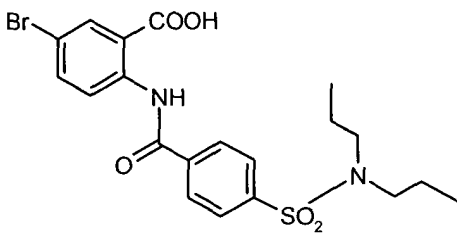
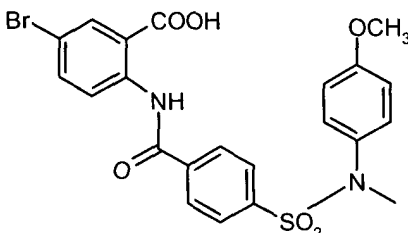
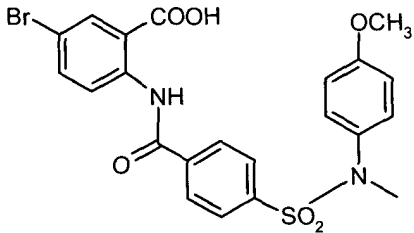
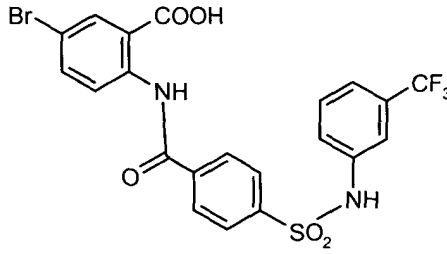
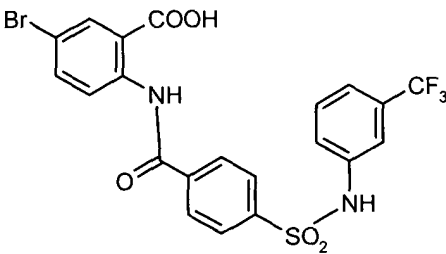
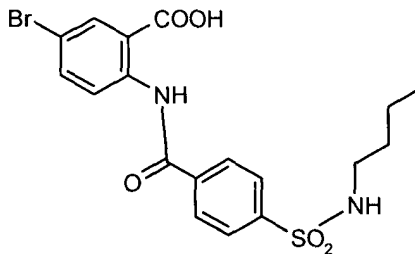
**3-Iodobenzoyl Chloride.** To a suspension of *m*-iodobenzoic acid (**21**, 5.0g, 20.1 mmol) suspended in CH<sub>2</sub>Cl<sub>2</sub> (60 mL) was added DMF (2 drops), followed by oxalyl chloride (20.1 mL of a 2 M solution in CH<sub>2</sub>Cl<sub>2</sub>, 40.2 mmol) under nitrogen atmosphere. After stirring for 18 h, the reaction mixture was nearly homogeneous.

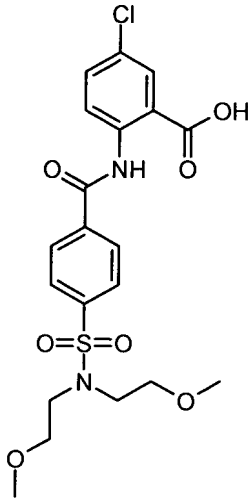
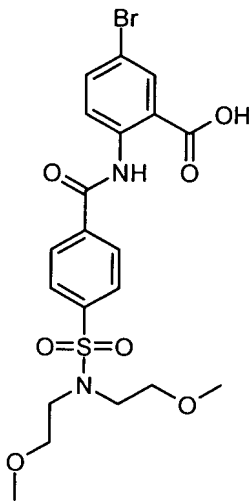
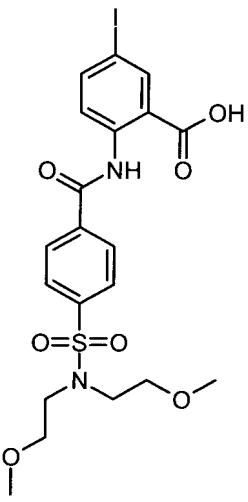
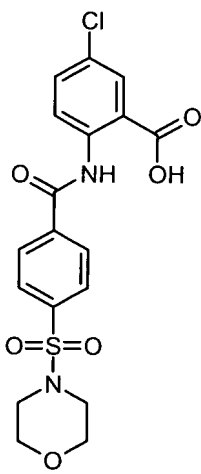
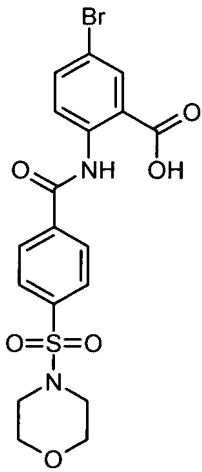
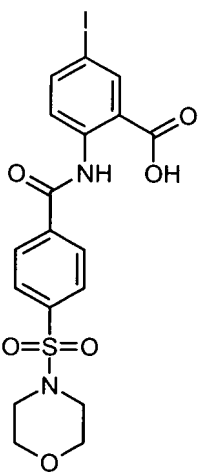


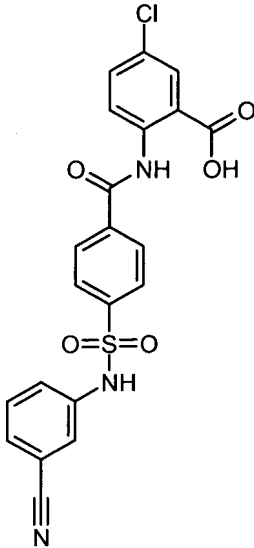
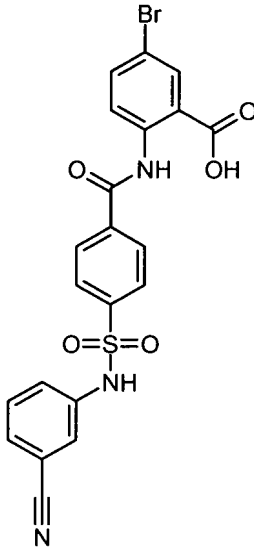
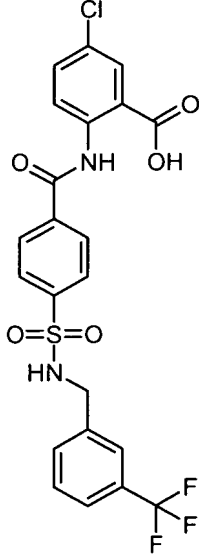
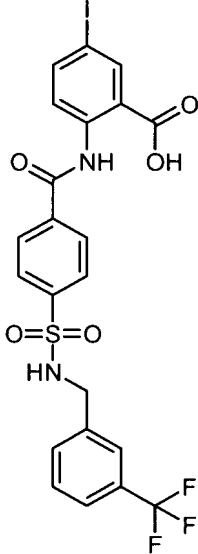
Acid chloride **22** was then concentrated, azeotroped with toluene (2 x 25 mL), and placed on a high vacuum.

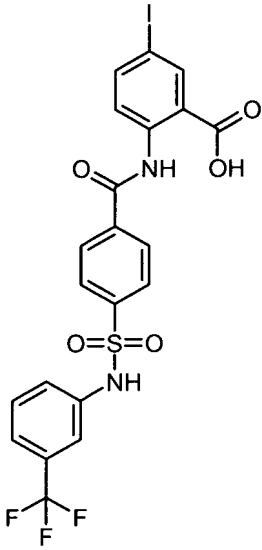
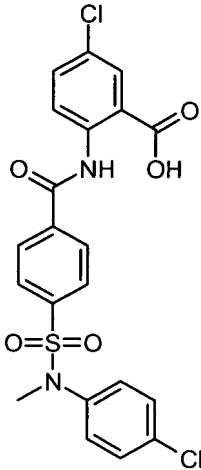
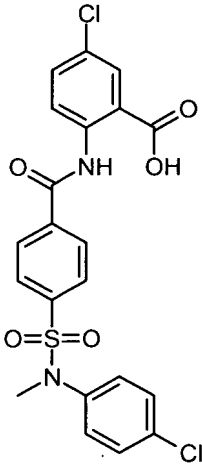
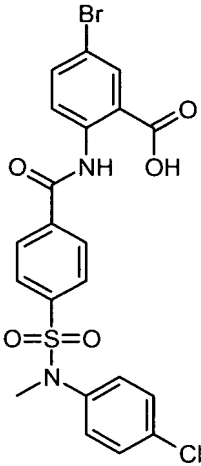
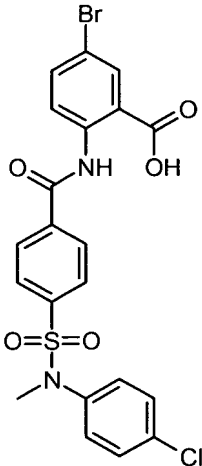
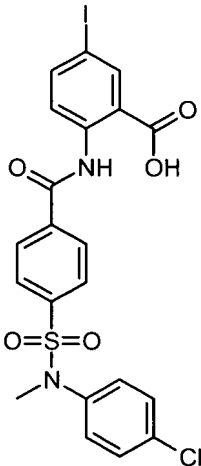
**Example 10: Additional Compounds Useful For Sterilization, Sanitation, Antisepsis, and Disinfection**

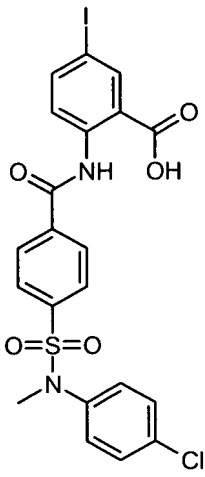
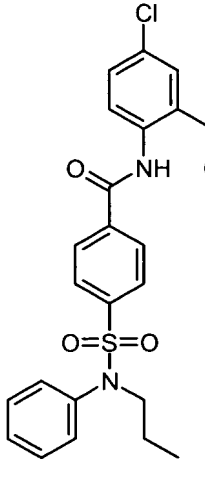
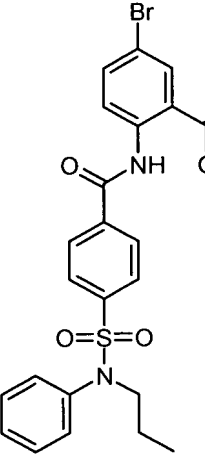
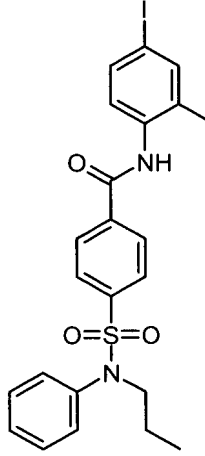
The following compounds may be synthesized using the methodology described above or via methods known in the art.

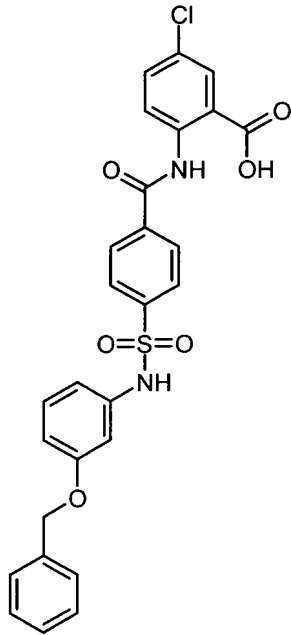
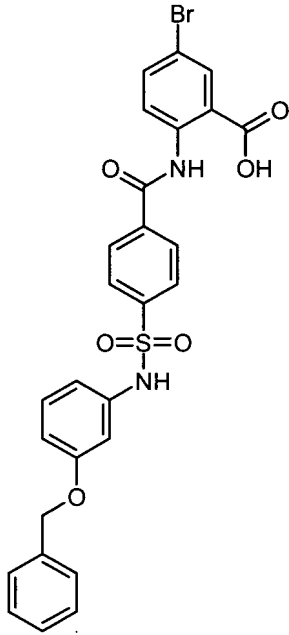
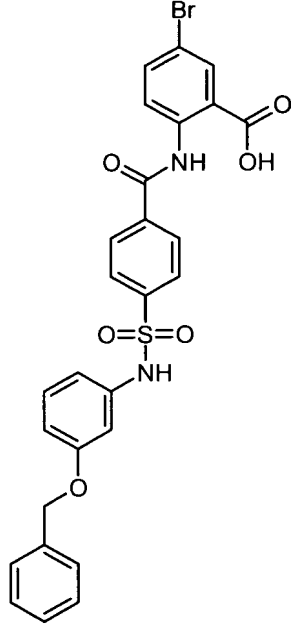
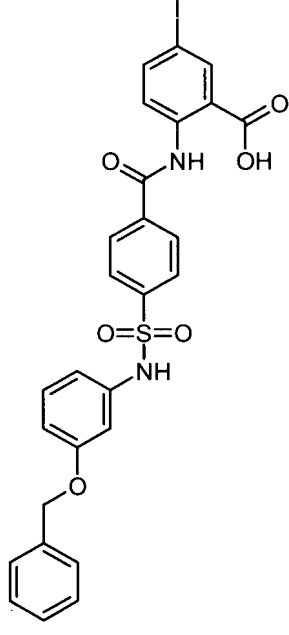
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<p>L-140209A</p> 	<p>L-140229</p> 
<p>L-140229A</p> 	<p>L-140240</p> 

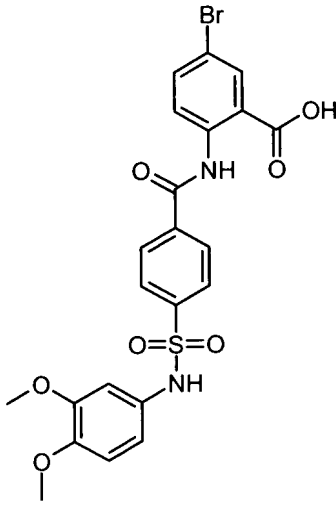
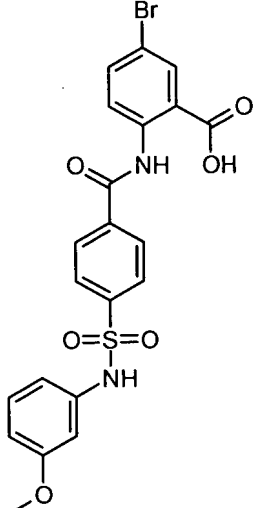
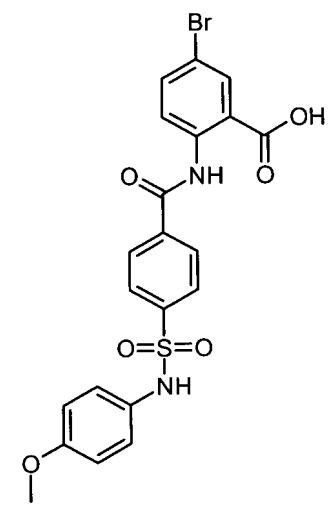
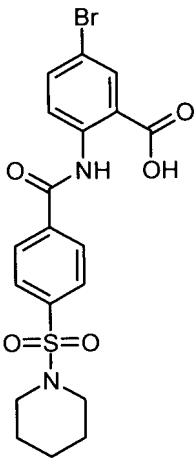
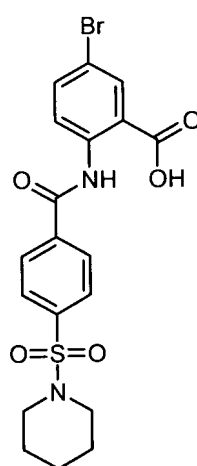
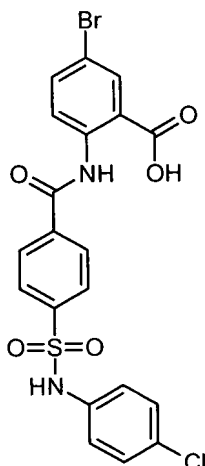
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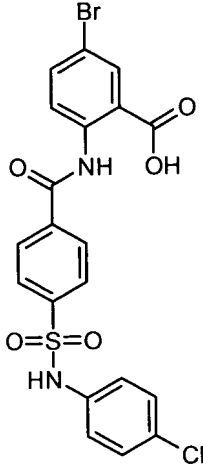
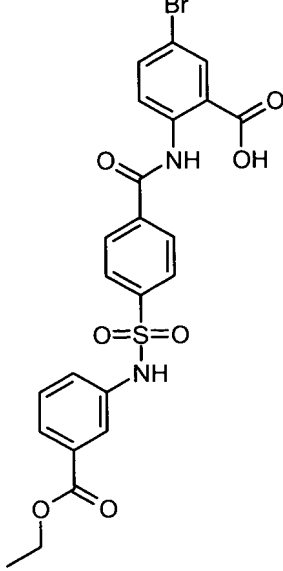
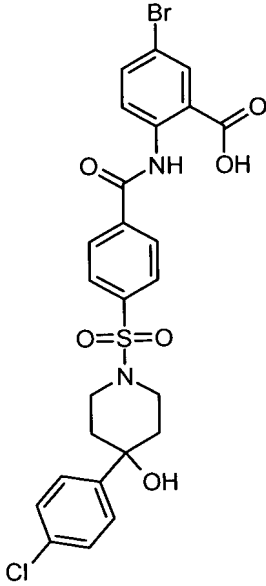
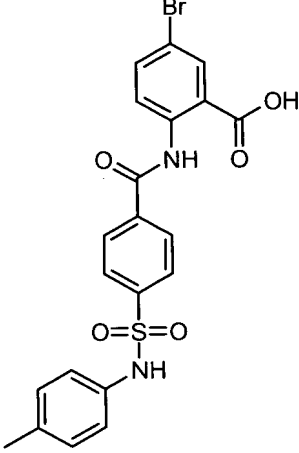
Compound No., Structure	Compound No., Structure
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<p data-bbox="277 825 407 856">L-159131</p>  <chem data-bbox="446 884 643 1434">Clc1ccc(NC(=O)c2ccc(S(=O)(=O)NCCc3ccc(C(F)(F)F)cc3)cc2)c(=O)O1</chem>	<p data-bbox="844 825 974 856">L-159133</p>  <chem data-bbox="1015 884 1211 1434">Ic1ccc(NC(=O)c2ccc(S(=O)(=O)NCCc3ccc(C(F)(F)F)cc3)cc2)c(=O)O1</chem>

Compound No., Structure	Compound No., Structure
<p data-bbox="277 180 407 212">L-159139</p> 	<p data-bbox="844 180 974 212">L-159157</p> 
<p data-bbox="277 825 428 856">L-159157A</p> 	<p data-bbox="844 825 974 856">L-159158</p> 
<p data-bbox="277 1381 428 1413">L-159158A</p> 	<p data-bbox="844 1381 974 1413">L-159161</p> 

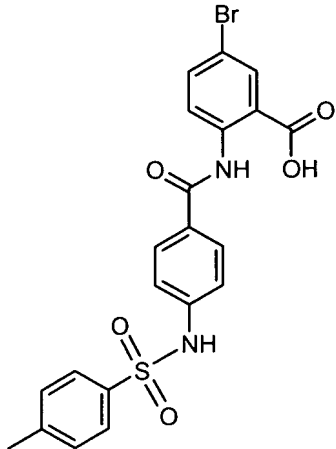
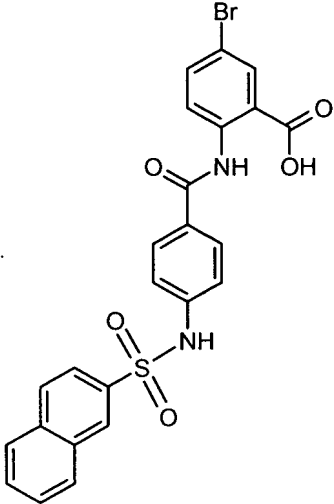
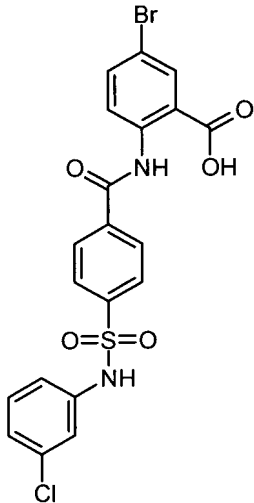
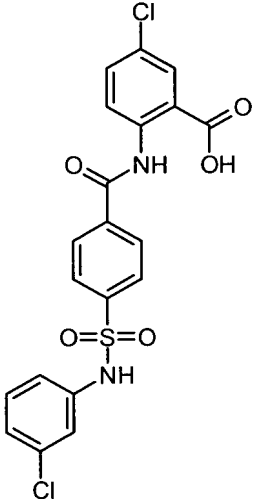
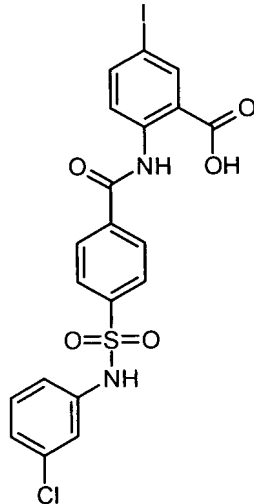
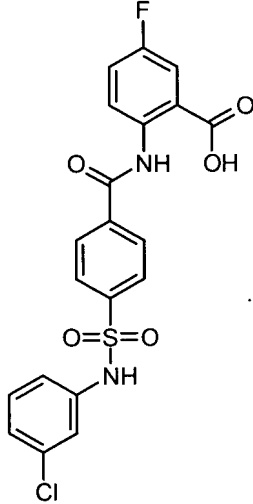
Compound No., Structure	Compound No., Structure
<p data-bbox="277 186 428 218">L-159161A</p>  <chem data-bbox="446 241 649 714">Clc1ccc(cc1)N(S(=O)(=O)c2ccc(cc2)NC(=O)Nc3cc(I)ccc3C(=O)O)c4ccccc4</chem>	<p data-bbox="841 186 976 218">L-159164</p>  <chem data-bbox="990 241 1193 714">Clc1ccc(cc1)N(S(=O)(=O)c2ccc(cc2)NC(=O)Nc3cc(Cl)ccc3C(=O)O)c4ccccc4</chem>
<p data-bbox="277 753 412 785">L-159165</p>  <chem data-bbox="430 808 633 1260">CCc1cccc(c1)N(S(=O)(=O)c2ccc(cc2)NC(=O)Nc3cc(Br)ccc3C(=O)O)c4ccccc4</chem>	<p data-bbox="841 753 976 785">L-159168</p>  <chem data-bbox="990 808 1193 1260">CCc1cccc(c1)N(S(=O)(=O)c2ccc(cc2)NC(=O)Nc3cc(I)ccc3C(=O)O)c4ccccc4</chem>

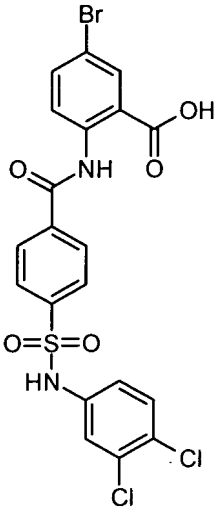
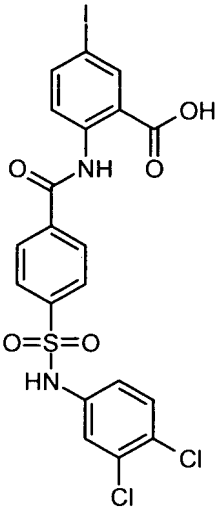
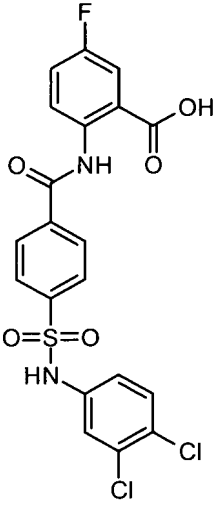
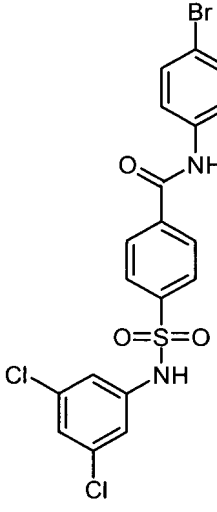
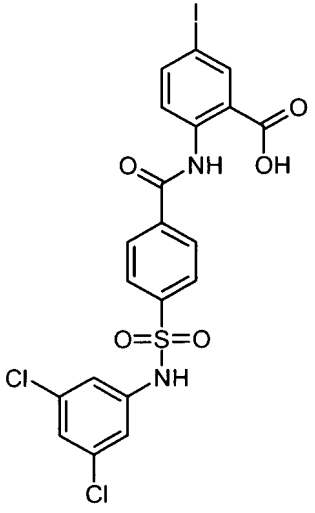
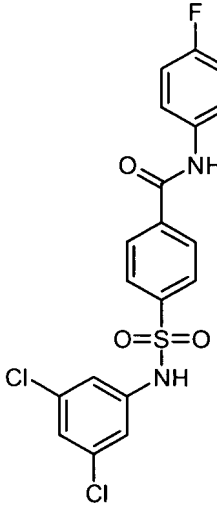
Compound No., Structure	Compound No., Structure
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<p data-bbox="272 913 423 947">L-159172A</p> 	<p data-bbox="841 913 971 947">L-159176</p> 

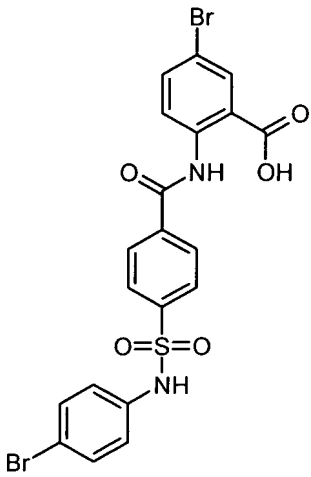
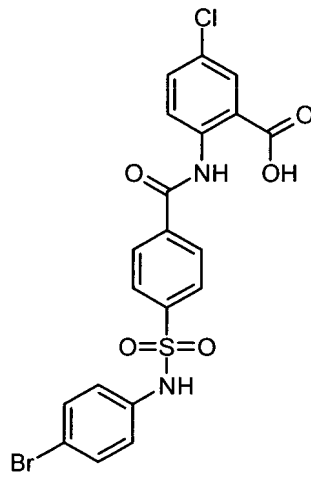
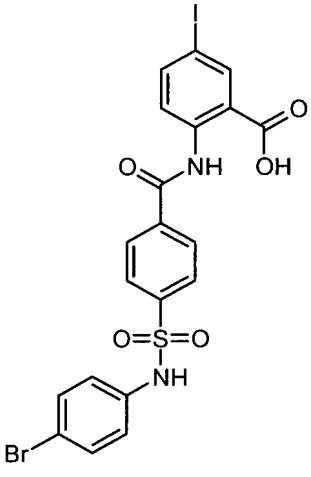
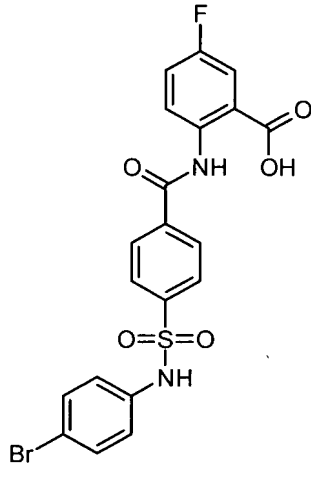
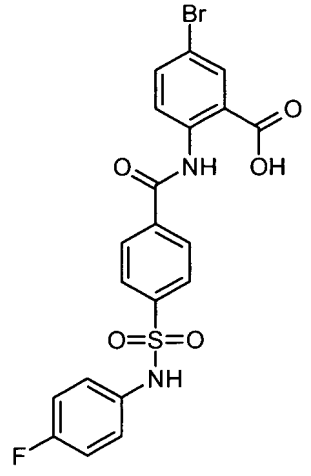
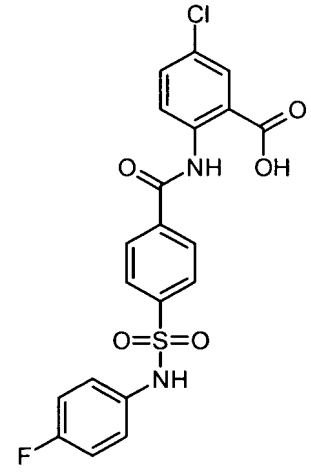
Compound No., Structure	Compound No., Structure
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<p data-bbox="267 802 397 837">L-170160</p> 	<p data-bbox="831 802 961 837">L-170166</p> 
<p data-bbox="267 1390 430 1425">L-170166A</p> 	<p data-bbox="831 1390 961 1425">L-170178</p> 

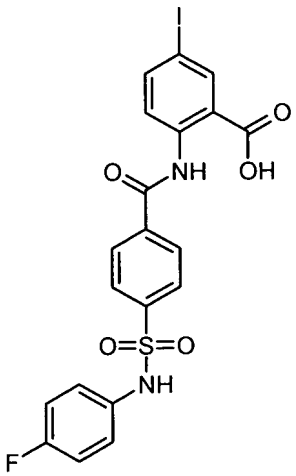
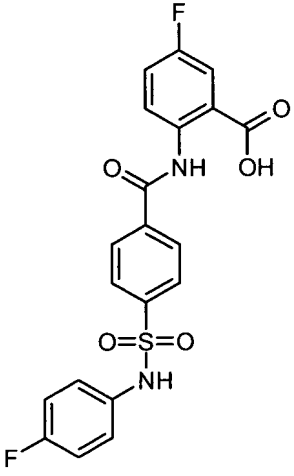
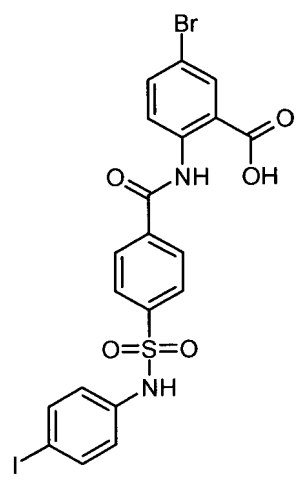
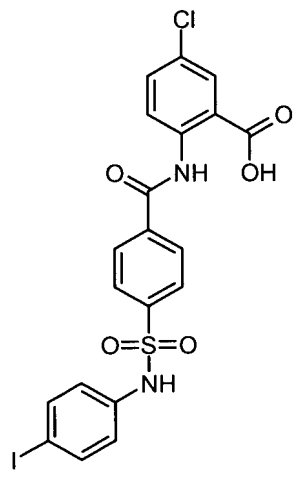
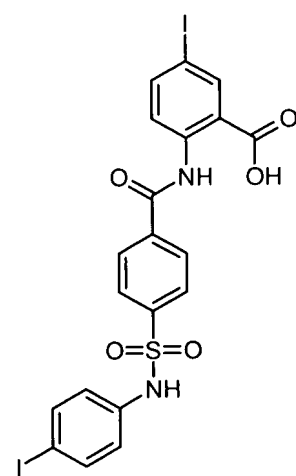
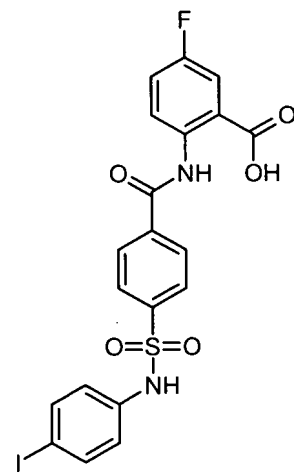
Compound No., Structure	Compound No., Structure
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<p data-bbox="266 863 399 894">L-170190</p> 	<p data-bbox="829 863 963 894">L-170196</p> 

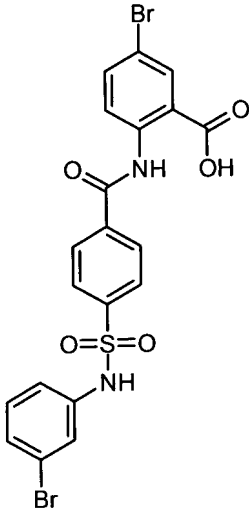
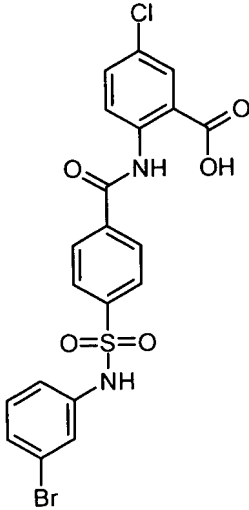
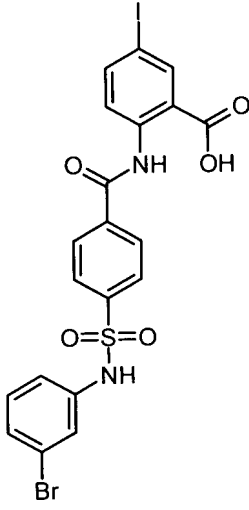
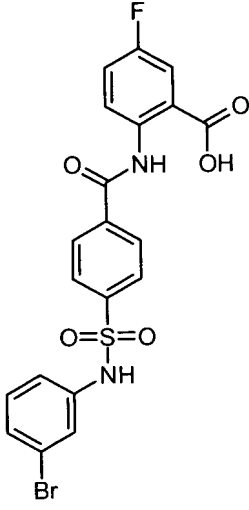
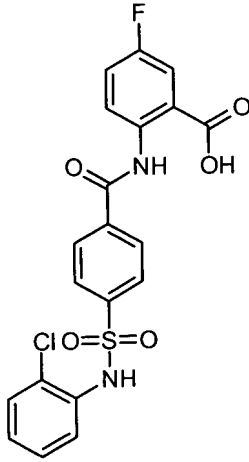
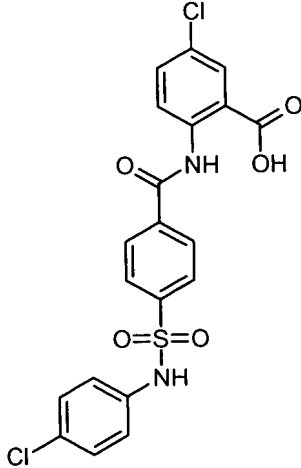


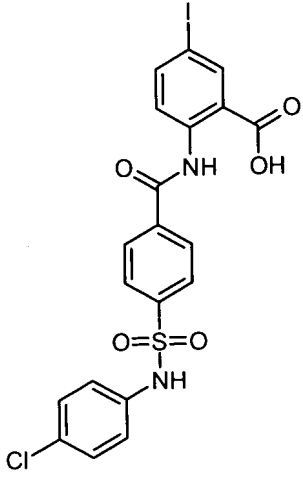
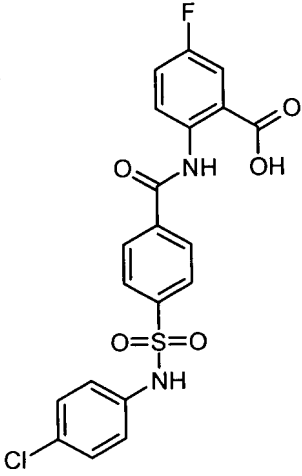
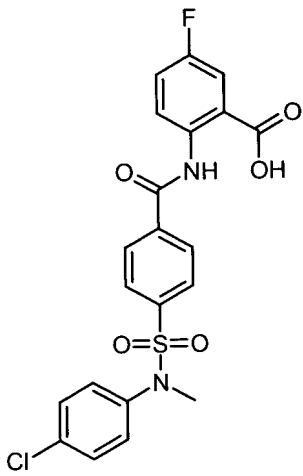
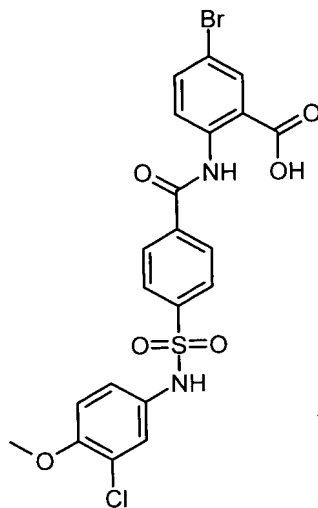
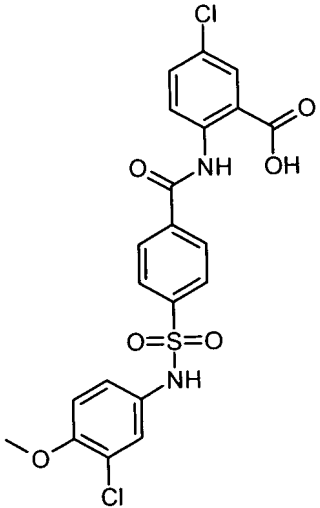
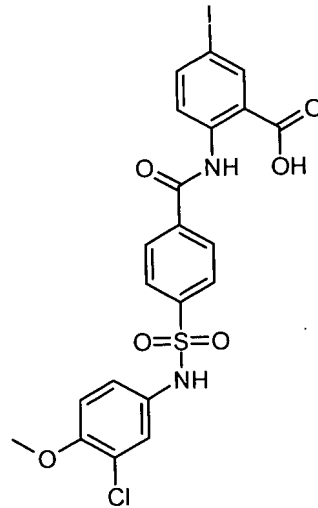
Compound No., Structure	Compound No., Structure
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<p data-bbox="271 785 406 816">L-181367</p> 	<p data-bbox="837 785 972 816">L-181368</p> 
<p data-bbox="271 1381 406 1413">L-181370</p> 	<p data-bbox="837 1381 972 1413">L-181371</p> 

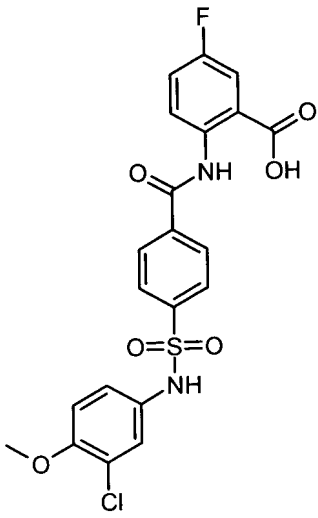
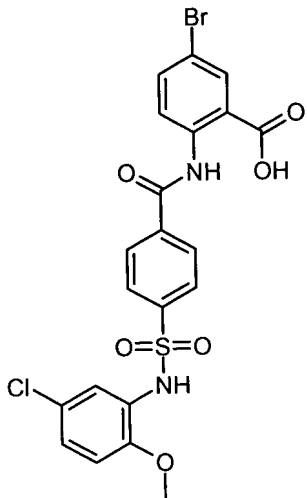
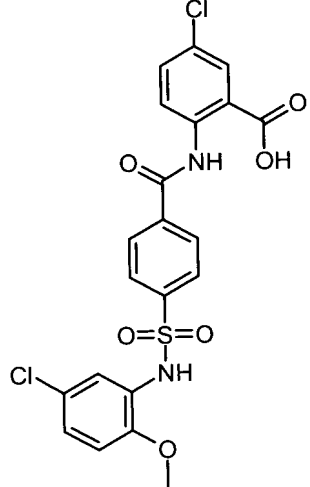
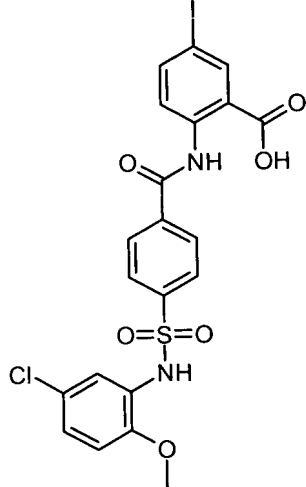
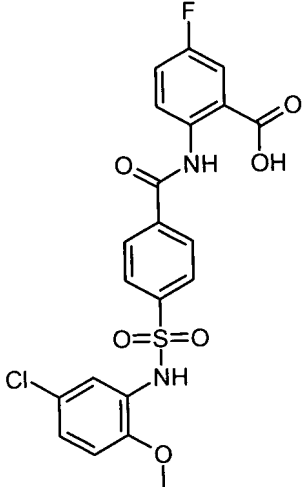
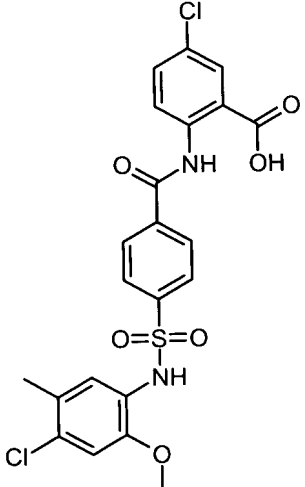
Compound No., Structure	Compound No., Structure
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<p data-bbox="266 770 402 800">L-181383</p>  <chem>O=C(O)c1ccc(NC(=O)c2ccc(S(=O)(=O)Nc3cc(Cl)cc(Cl)c3)cc2)cc1F</chem>	<p data-bbox="834 770 971 800">L-181385</p>  <chem>O=C(O)c1ccc(NC(=O)c2ccc(S(=O)(=O)Nc3cc(Cl)cc(Cl)c3)cc2)cc1Br</chem>
<p data-bbox="266 1365 402 1394">L-181388</p>  <chem>O=C(O)c1ccc(NC(=O)c2ccc(S(=O)(=O)Nc3cc(Cl)cc(Cl)c3)cc2)cc1I</chem>	<p data-bbox="834 1365 971 1394">L-181389</p>  <chem>O=C(O)c1ccc(NC(=O)c2ccc(S(=O)(=O)Nc3cc(Cl)cc(Cl)c3)cc2)cc1F</chem>

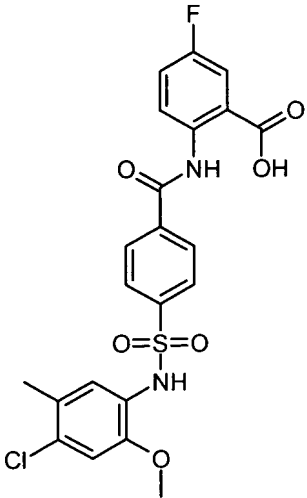
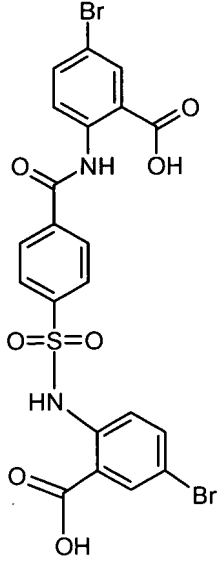
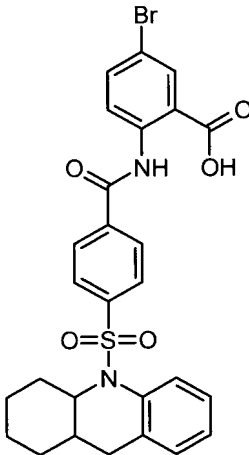
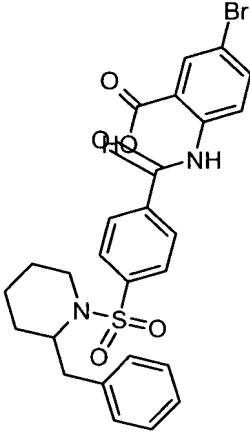
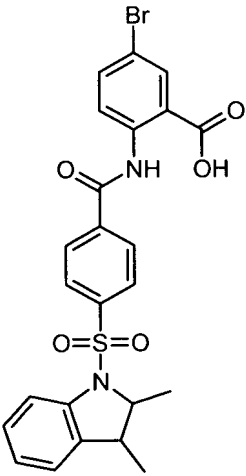
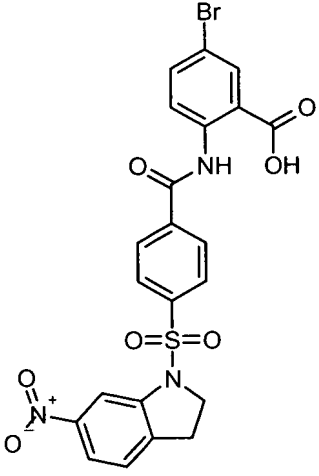
Compound No., Structure	Compound No., Structure
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<p data-bbox="269 741 402 772">L-181394</p> 	<p data-bbox="837 741 971 772">L-181395</p> 
<p data-bbox="269 1297 402 1329">L-181397</p> 	<p data-bbox="837 1297 971 1329">L-181398</p> 

Compound No., Structure	Compound No., Structure
<p data-bbox="267 174 397 210">L-181400</p>  <chem data-bbox="397 231 690 703">O=C(O)c1cc(NC(=O)c2ccc(S(=O)(=O)Nc3ccc(F)cc3)cc2)cc(I)c1</chem>	<p data-bbox="836 174 966 210">L-181401</p>  <chem data-bbox="966 231 1258 703">O=C(O)c1cc(NC(=O)c2ccc(S(=O)(=O)Nc3ccc(F)cc3)cc2)cc(F)c1</chem>
<p data-bbox="267 741 397 777">L-181403</p>  <chem data-bbox="397 787 690 1260">O=C(O)c1cc(NC(=O)c2ccc(S(=O)(=O)Nc3ccc(I)cc3)cc2)cc(Br)c1</chem>	<p data-bbox="836 741 966 777">L-181404</p>  <chem data-bbox="966 787 1258 1260">O=C(O)c1cc(NC(=O)c2ccc(S(=O)(=O)Nc3ccc(I)cc3)cc2)cc(Cl)c1</chem>
<p data-bbox="267 1297 397 1333">L-181406</p>  <chem data-bbox="397 1344 690 1816">O=C(O)c1cc(NC(=O)c2ccc(S(=O)(=O)Nc3ccc(I)cc3)cc2)cc(I)c1</chem>	<p data-bbox="836 1297 966 1333">L-181407</p>  <chem data-bbox="966 1344 1258 1816">O=C(O)c1cc(NC(=O)c2ccc(S(=O)(=O)Nc3ccc(I)cc3)cc2)cc(F)c1</chem>

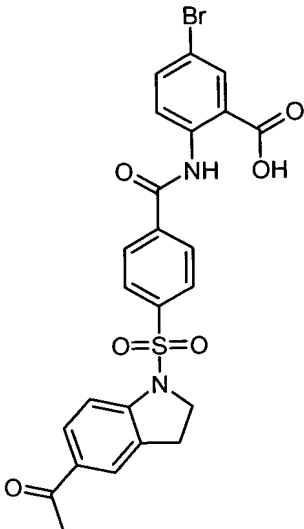
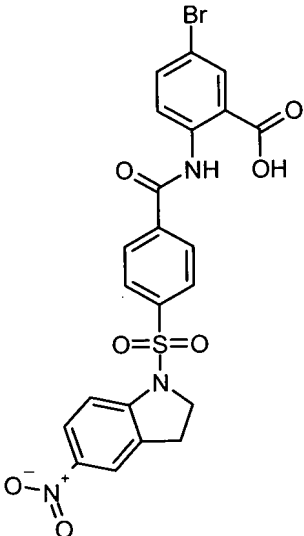
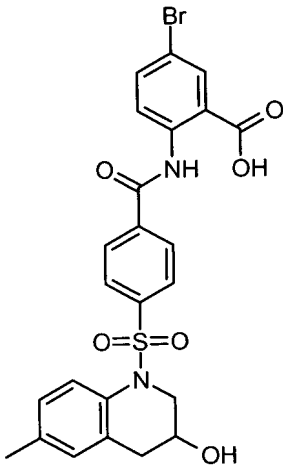
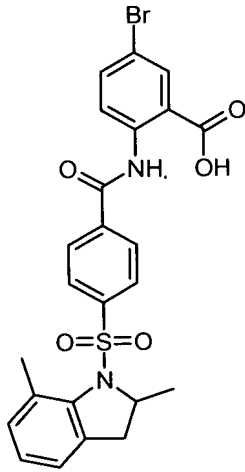
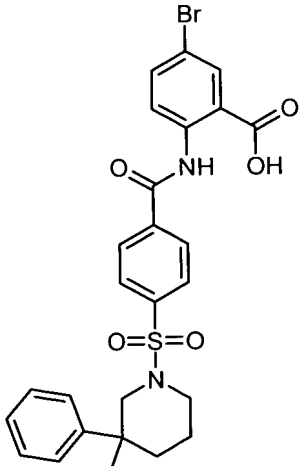
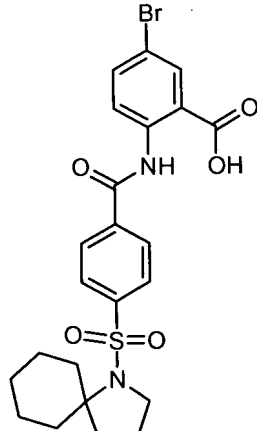
Compound No., Structure	Compound No., Structure
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<p data-bbox="261 785 391 816">L-181412</p> 	<p data-bbox="829 785 959 816">L-181413</p> 
<p data-bbox="261 1390 391 1421">L-181419</p> 	<p data-bbox="829 1390 959 1421">L-181421</p> 

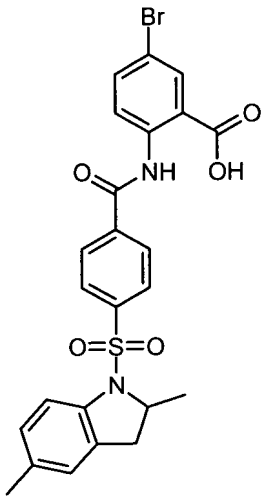
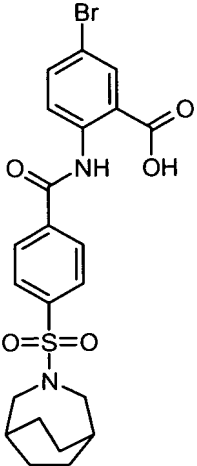
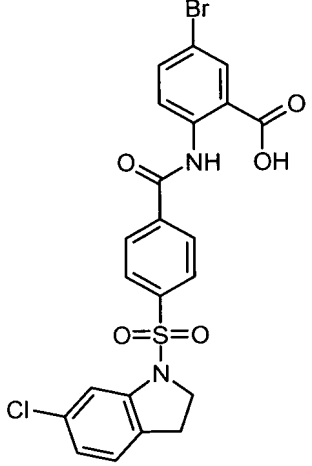
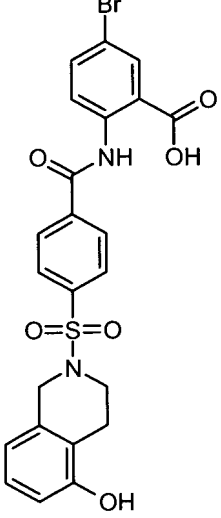
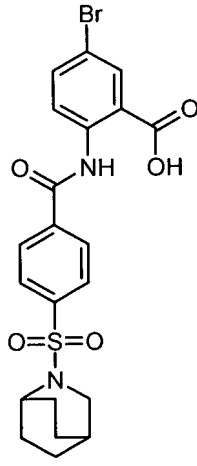
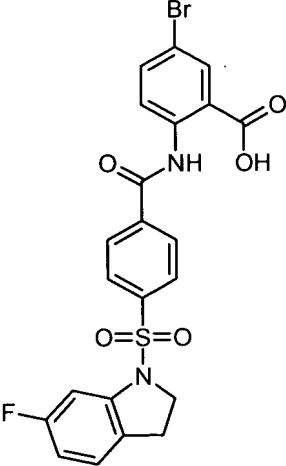
Compound No., Structure	Compound No., Structure
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<p data-bbox="256 745 397 787">L-181427</p>  <chem data-bbox="389 798 690 1270">Fc1ccc(NC(=O)S(=O)(=O)N(C)C(=O)c2ccc(Cl)cc2)cc1</chem>	<p data-bbox="820 745 966 787">L-181429</p>  <chem data-bbox="950 787 1266 1291">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)NC(=O)c3cc(F)cc(Br)c3</chem>
<p data-bbox="256 1333 406 1375">L-181430</p>  <chem data-bbox="397 1396 714 1900">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)NC(=O)c3cc(F)cc(Cl)c3</chem>	<p data-bbox="820 1333 966 1375">L-181432</p>  <chem data-bbox="958 1386 1274 1890">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)NC(=O)c3cc(F)cc(I)c3</chem>

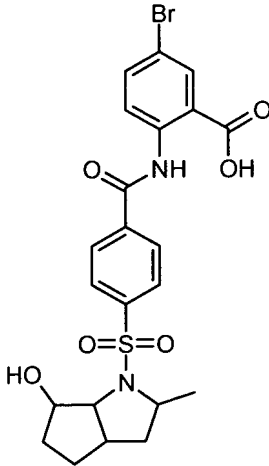
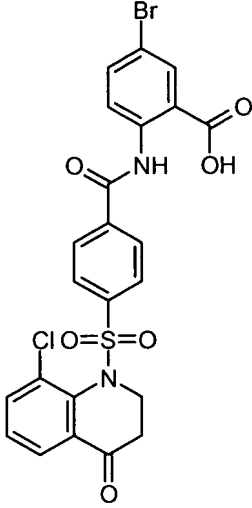
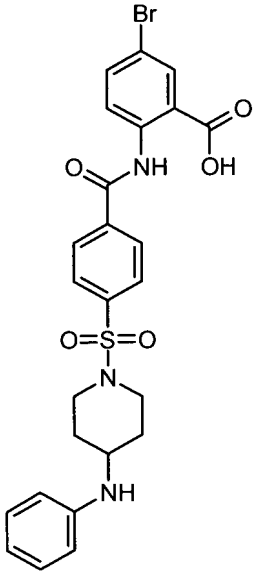
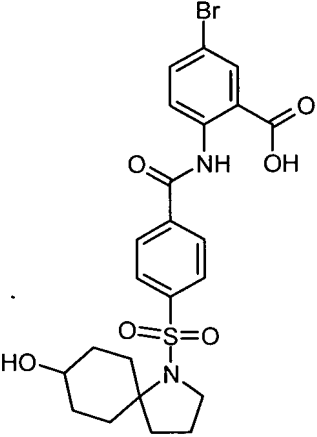
Compound No., Structure	Compound No., Structure
<p data-bbox="248 184 378 216">L-181433</p>  <chem data-bbox="370 237 686 751">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)C(=O)Nc3ccc(F)cc3C(=O)O</chem>	<p data-bbox="815 174 945 205">L-181435</p>  <chem data-bbox="937 226 1242 720">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)C(=O)Nc3ccc(Br)cc3C(=O)O</chem>
<p data-bbox="256 783 386 814">L-181436</p>  <chem data-bbox="378 846 686 1329">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)C(=O)Nc3cc(Cl)ccc3C(=O)O</chem>	<p data-bbox="824 783 954 814">L-181438</p>  <chem data-bbox="946 835 1250 1318">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)C(=O)Nc3cc(I)ccc3C(=O)O</chem>
<p data-bbox="264 1371 394 1402">L-181439</p>  <chem data-bbox="394 1434 698 1917">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)C(=O)Nc3ccc(F)cc3C(=O)O</chem>	<p data-bbox="833 1371 963 1402">L-181442</p>  <chem data-bbox="963 1423 1266 1906">COc1cc(Cl)ccc1NS(=O)(=O)c2ccc(cc2)C(=O)Nc3cc(Cl)ccc3C(=O)O</chem>

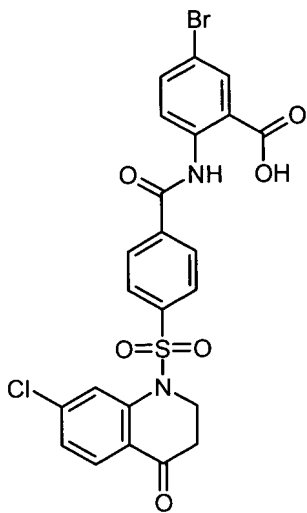
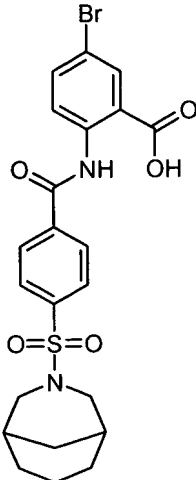
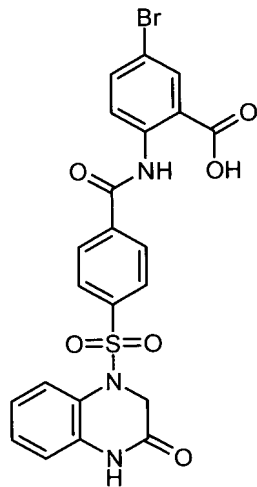
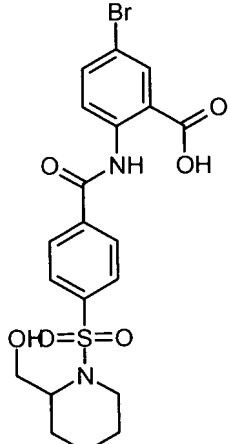
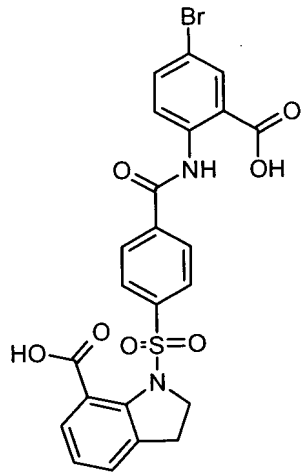
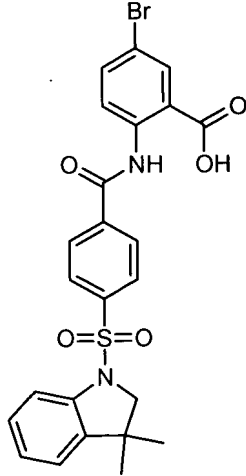
Compound No., Structure	Compound No., Structure
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<p>L-199155</p> 	<p>L-199156</p> 
<p>L-199157</p> 	<p>L-199158</p> 

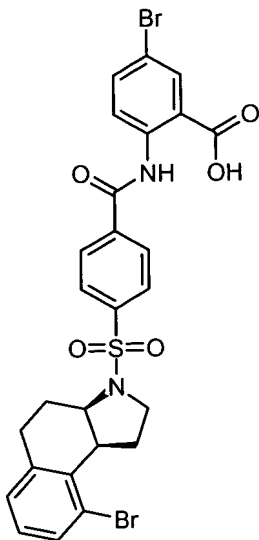
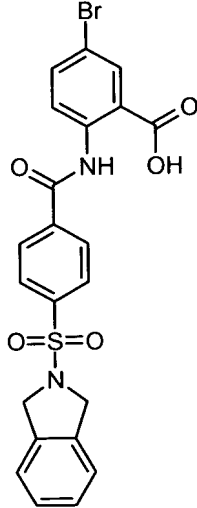
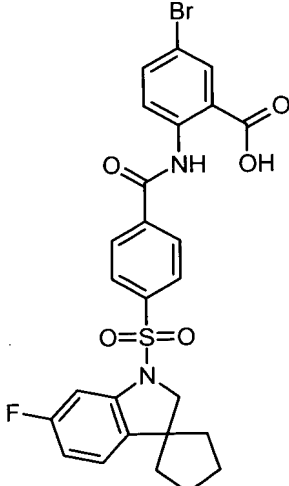
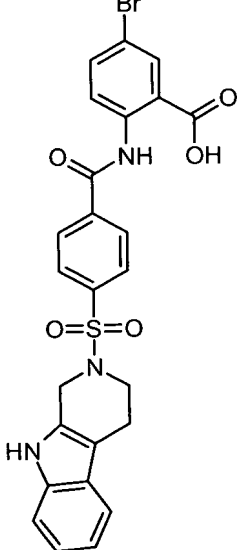


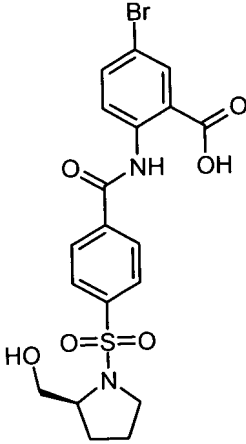
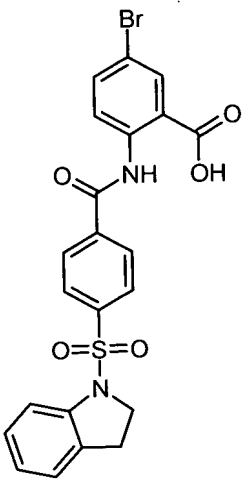
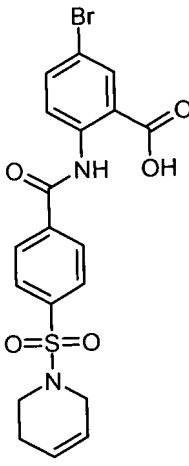
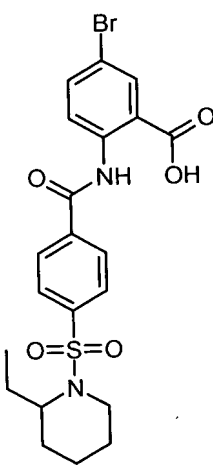
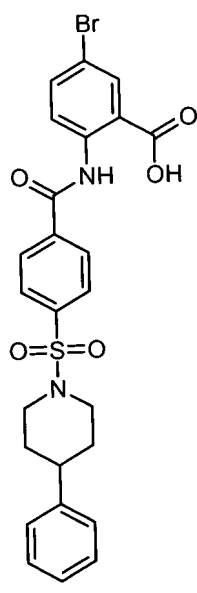
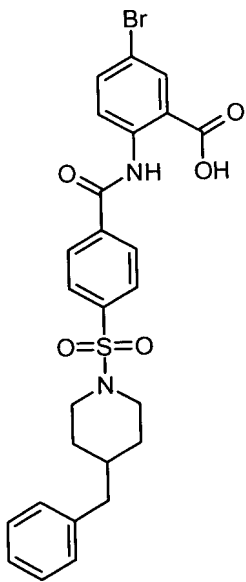
Compound No., Structure	Compound No., Structure
<p>L-199159</p> 	<p>L-199160</p> 
<p>L-199161</p> 	<p>L-199162</p> 
<p>L-199163</p> 	<p>L-199164</p> 

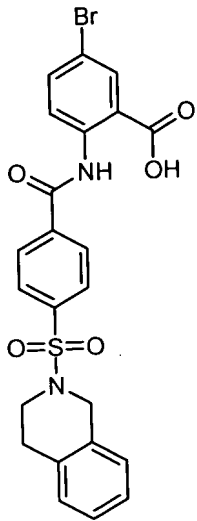
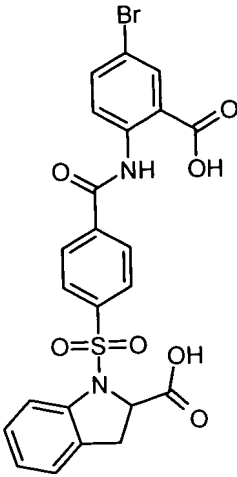
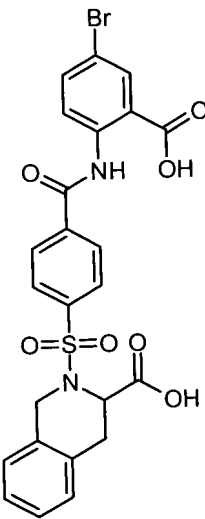
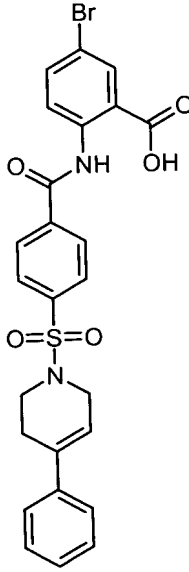
Compound No., Structure	Compound No., Structure
<p data-bbox="269 170 402 201">L-199165</p> 	<p data-bbox="837 170 971 201">L-199166</p> 
<p data-bbox="269 762 402 793">L-199167</p> 	<p data-bbox="837 762 971 793">L-199168</p> 
<p data-bbox="269 1377 402 1409">L-199169</p> 	<p data-bbox="837 1377 971 1409">L-199170</p> 

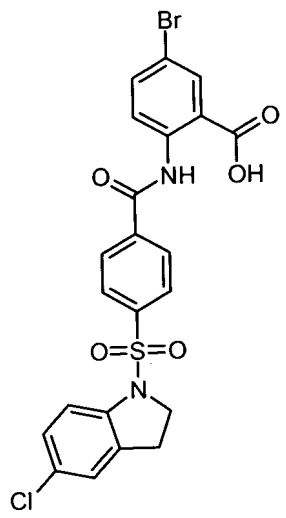
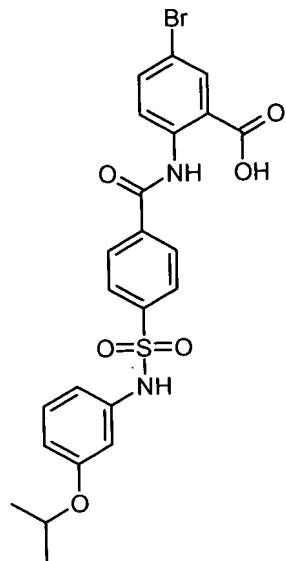
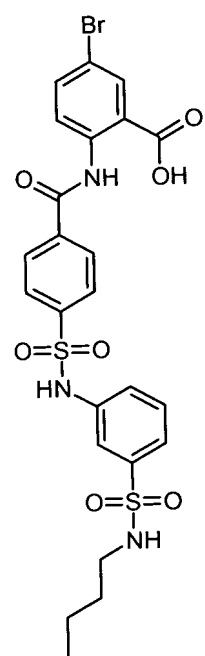
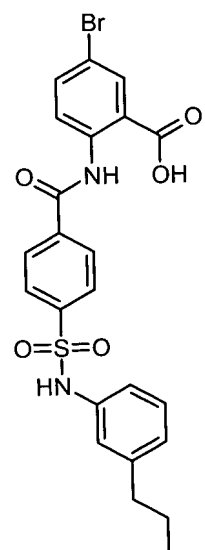
Compound No., Structure	Compound No., Structure
<p data-bbox="277 170 402 197">L-199171</p>  <p>The structure of L-199171 consists of a 3-bromo-4-((4-((1-methyl-2-hydroxy-2,3-dihydro-1H-indeno[1,2-b]pyridin-5-yl)sulfonyl)phenyl)carbamoyl)benzoic acid moiety. It features a benzene ring with a bromine atom at the 3-position and a carboxylic acid group at the 1-position. An amide linkage connects the 4-position to a para-substituted phenyl ring, which is further linked via a sulfonyl group to a 1-methyl-2-hydroxy-2,3-dihydro-1H-indeno[1,2-b]pyridin-5-yl group.</p>	<p data-bbox="842 170 967 197">L-199172</p>  <p>The structure of L-199172 consists of a 3-bromo-4-((4-((8-chloro-1,2,3,4-tetrahydroquinolin-2(1H)-one-5-yl)sulfonyl)phenyl)carbamoyl)benzoic acid moiety. It features a benzene ring with a bromine atom at the 3-position and a carboxylic acid group at the 1-position. An amide linkage connects the 4-position to a para-substituted phenyl ring, which is further linked via a sulfonyl group to an 8-chloro-1,2,3,4-tetrahydroquinolin-2(1H)-one-5-yl group.</p>
<p data-bbox="277 770 402 798">L-199173</p>  <p>The structure of L-199173 consists of a 3-bromo-4-((4-((4-phenylpiperidin-1-yl)sulfonyl)phenyl)carbamoyl)benzoic acid moiety. It features a benzene ring with a bromine atom at the 3-position and a carboxylic acid group at the 1-position. An amide linkage connects the 4-position to a para-substituted phenyl ring, which is further linked via a sulfonyl group to a 4-phenylpiperidin-1-yl group.</p>	<p data-bbox="842 770 967 798">L-199174</p>  <p>The structure of L-199174 consists of a 3-bromo-4-((4-((4-hydroxy-4,5,6,7-tetrahydroindeno[1,2-b]pyridin-5-yl)sulfonyl)phenyl)carbamoyl)benzoic acid moiety. It features a benzene ring with a bromine atom at the 3-position and a carboxylic acid group at the 1-position. An amide linkage connects the 4-position to a para-substituted phenyl ring, which is further linked via a sulfonyl group to a 4-hydroxy-4,5,6,7-tetrahydroindeno[1,2-b]pyridin-5-yl group.</p>

Compound No., Structure	Compound No., Structure
<p>L-199175</p> 	<p>L-199176</p> 
<p>L-199177</p> 	<p>L-199178</p> 
<p>L-199179</p> 	<p>L-199180</p> 

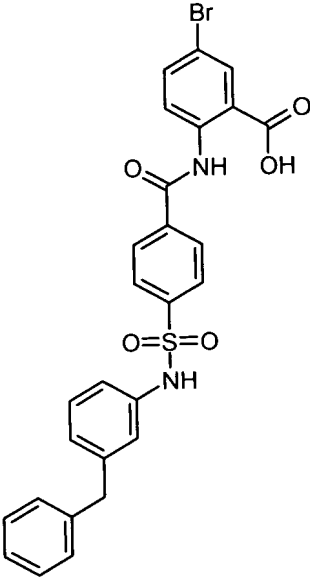
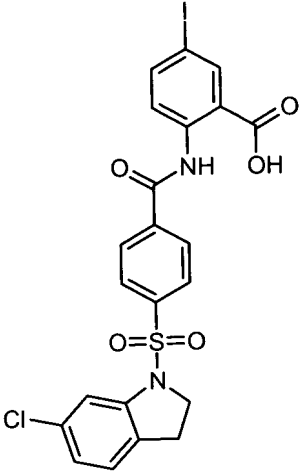
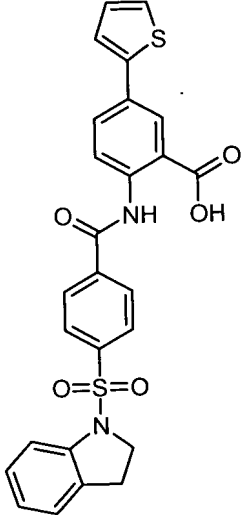
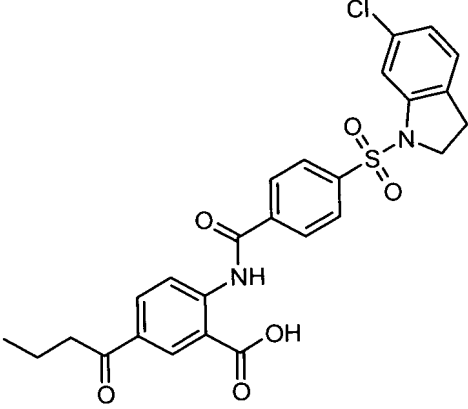
Compound No., Structure	Compound No., Structure
<p data-bbox="261 174 391 205">L-199181</p> 	<p data-bbox="837 174 967 205">L-199182</p> 
<p data-bbox="261 804 391 835">L-199183</p> 	<p data-bbox="837 804 967 835">L-199184</p> 

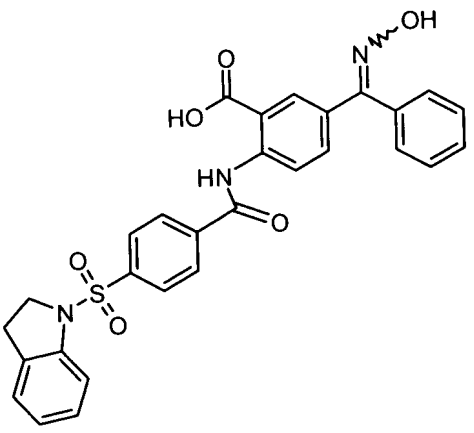
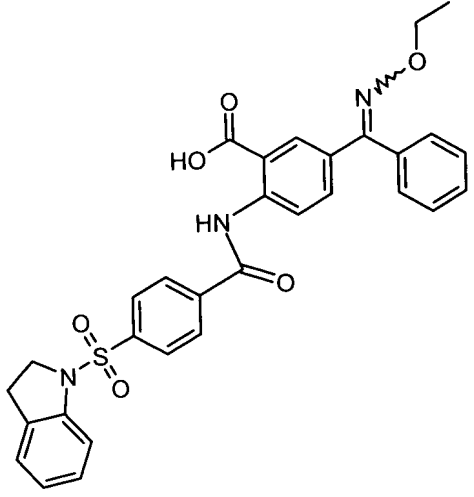
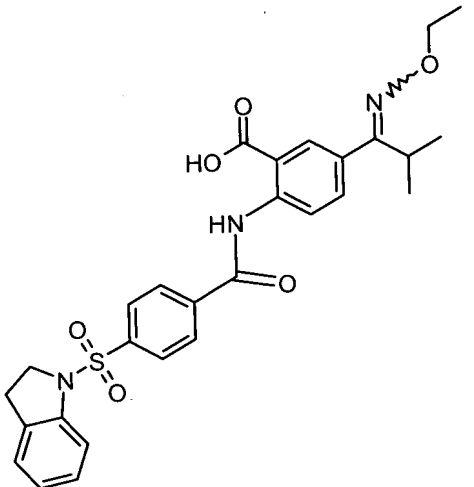
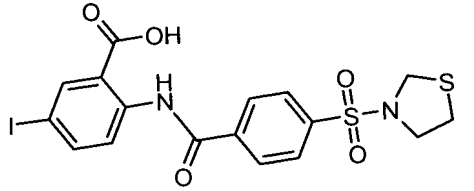
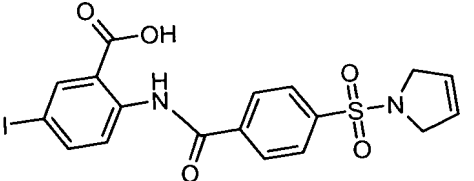
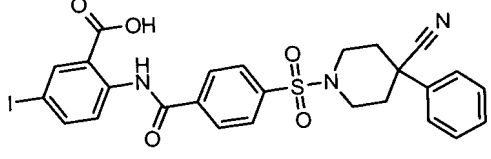
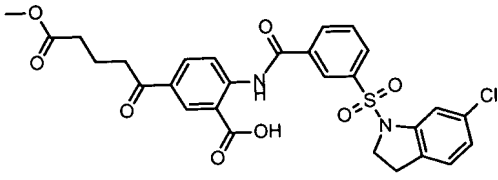
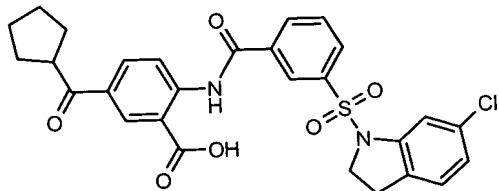
Compound No., Structure	Compound No., Structure
<p data-bbox="245 184 375 216">L-199185</p> 	<p data-bbox="813 174 943 205">L-199186</p> 
<p data-bbox="253 758 383 789">L-199187</p> 	<p data-bbox="821 747 951 779">L-199188</p> 
<p data-bbox="269 1314 399 1346">L-199189</p> 	<p data-bbox="837 1293 967 1325">L-199190</p> 

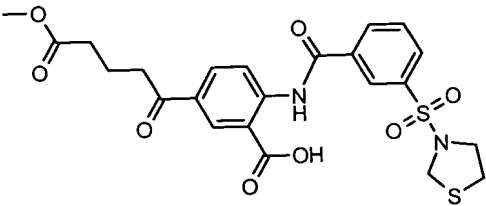
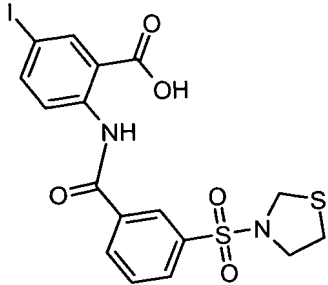
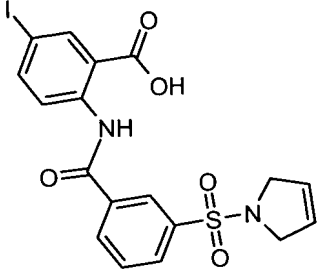
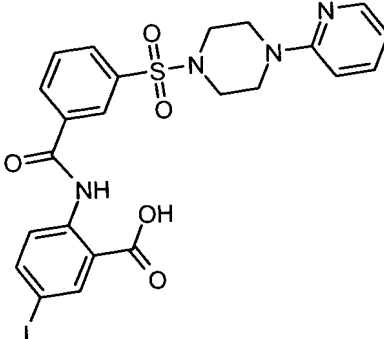
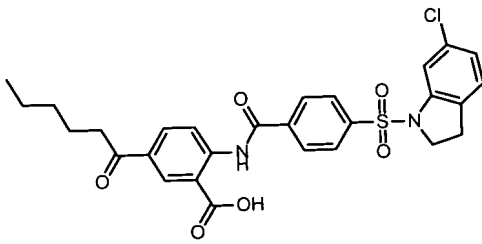
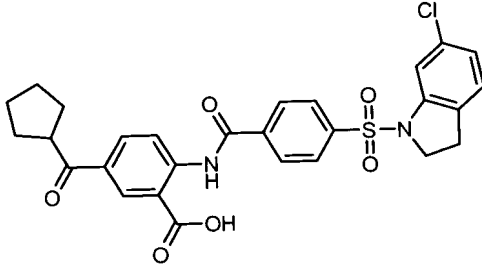
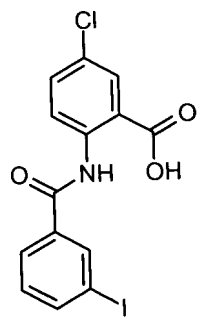
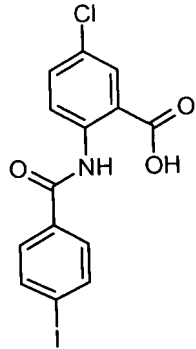
Compound No., Structure	Compound No., Structure
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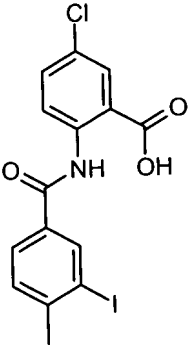
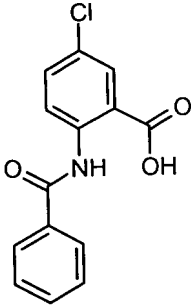
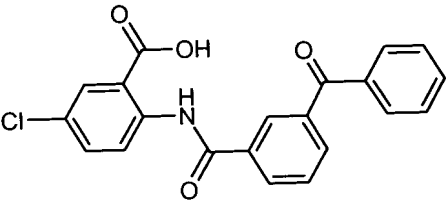
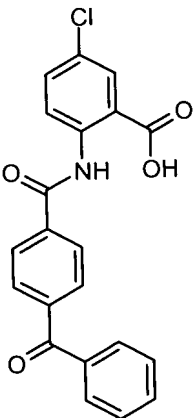
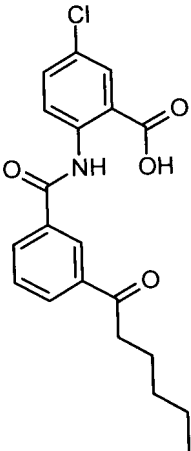
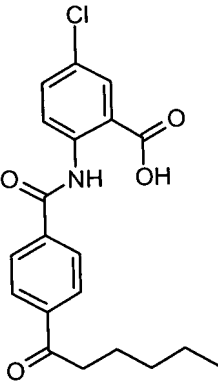
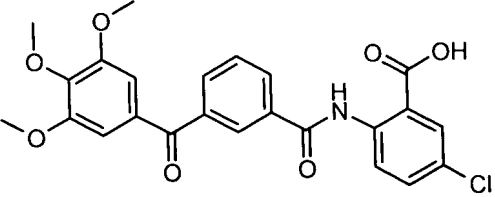
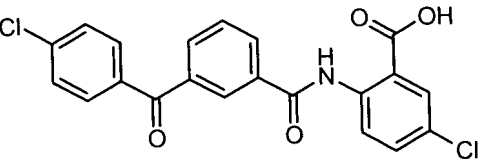
Compound No., Structure	Compound No., Structure
<p data-bbox="251 178 381 220">L-199195</p>  <chem data-bbox="381 231 665 745">BrC1=CC=C(C(=O)N1C(=O)c2ccc(cc2)S(=O)(=O)N3C4C=CC=C(C=C4)CN3Cl)C(=O)O</chem>	<p data-bbox="820 178 950 220">L-199196</p>  <chem data-bbox="950 220 1234 777">BrC1=CC=C(C(=O)N1C(=O)c2ccc(cc2)S(=O)(=O)Nc3ccc(cc3)OC(C)C)C(=O)O</chem>
<p data-bbox="251 835 381 877">L-199197</p>  <chem data-bbox="430 882 633 1533">BrC1=CC=C(C(=O)N1C(=O)c2ccc(cc2)S(=O)(=O)Nc3ccc(cc3)S(=O)(=O)NCCCC)C(=O)O</chem>	<p data-bbox="820 835 950 877">L-199198</p>  <chem data-bbox="998 871 1201 1417">BrC1=CC=C(C(=O)N1C(=O)c2ccc(cc2)S(=O)(=O)Nc3ccc(cc3)CCC)C(=O)O</chem>

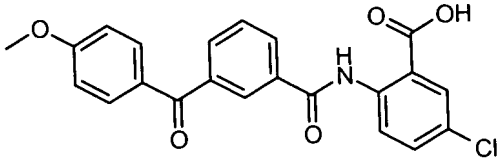
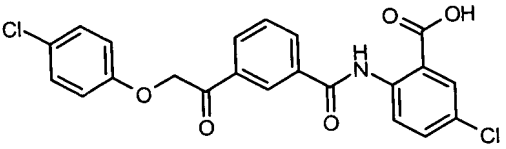
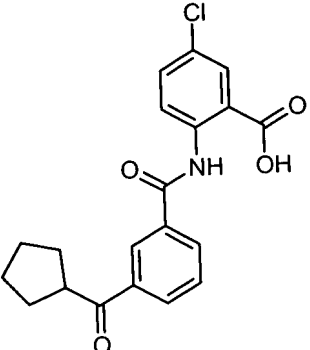
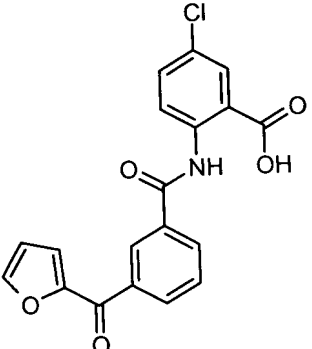
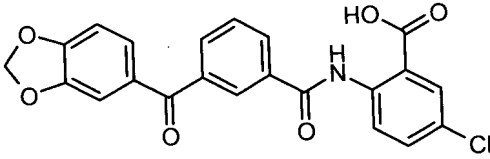
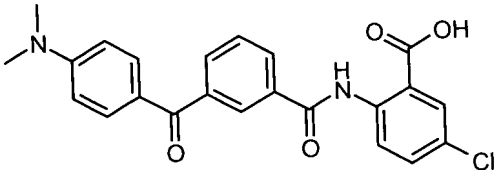
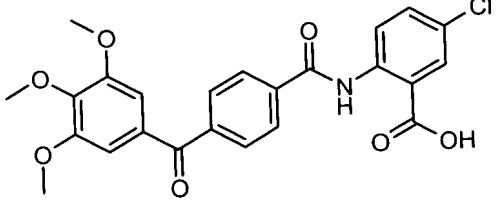
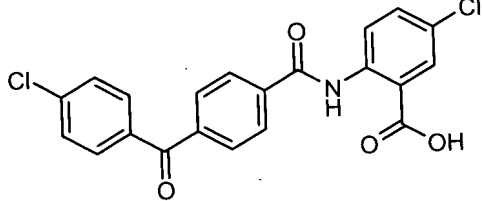
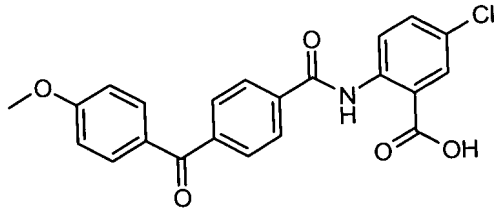
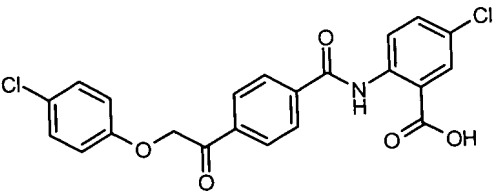


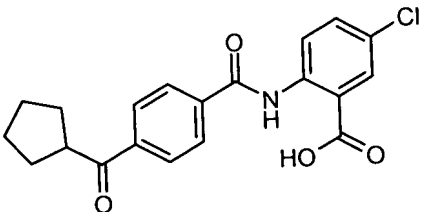
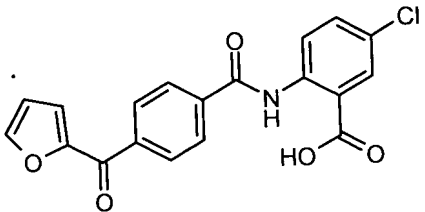
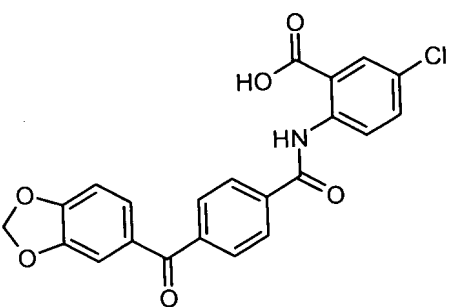
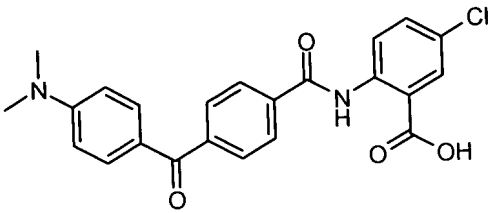
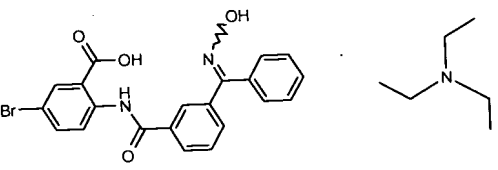
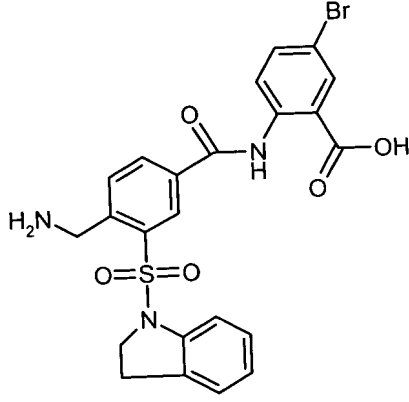
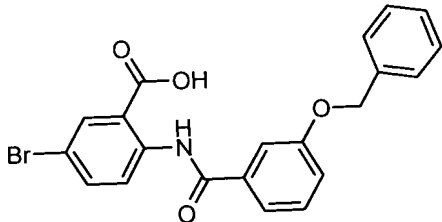
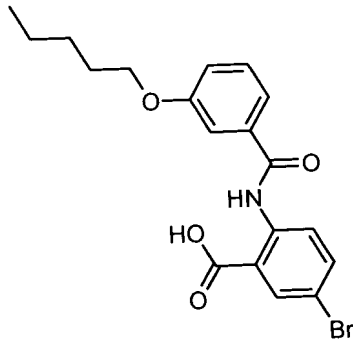
Compound No., Structure	Compound No., Structure
<p data-bbox="250 180 380 216">L-199199</p> 	<p data-bbox="816 170 946 205">L-217790</p> 
<p data-bbox="250 856 380 892">L-217791</p> 	<p data-bbox="816 846 946 882">L-218343</p> 

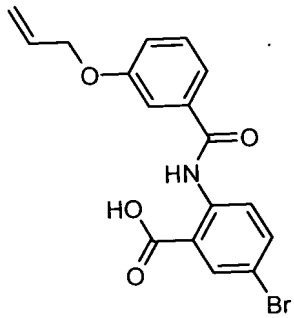
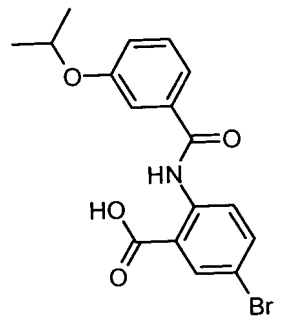
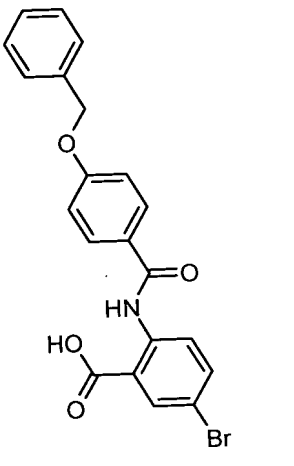
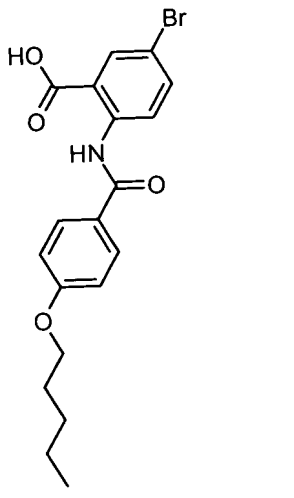
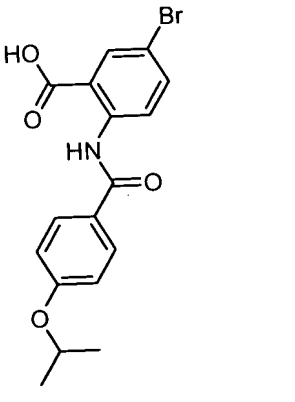
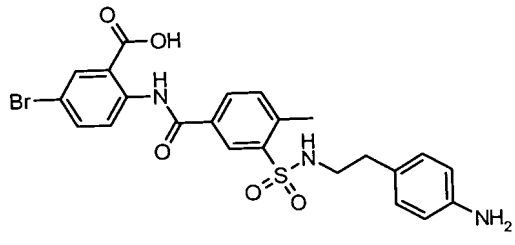
Compound No., Structure	Compound No., Structure
<p>L-502902</p> 	<p>L-502903</p> 
<p>L-502904</p> 	<p>PHA-500140</p> 
<p>PHA-500152</p> 	<p>PHA-500200</p> 
<p>PHA-500218</p> 	<p>PHA-500219</p> 

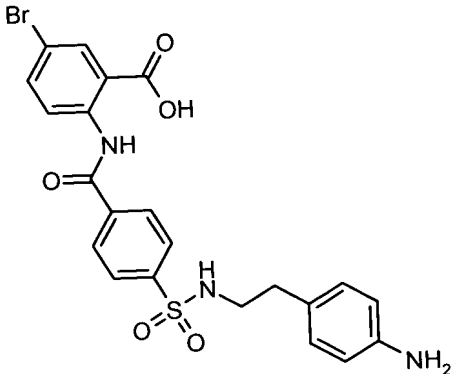
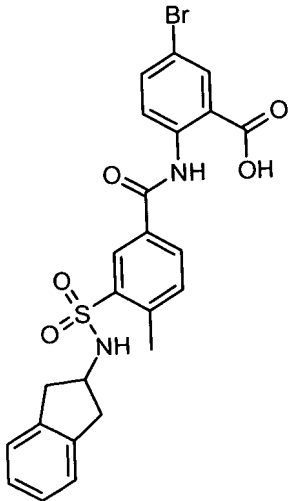
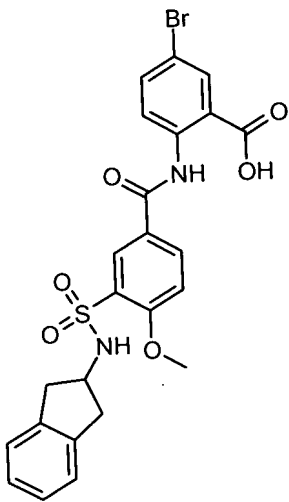
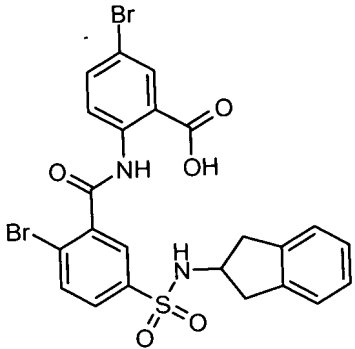
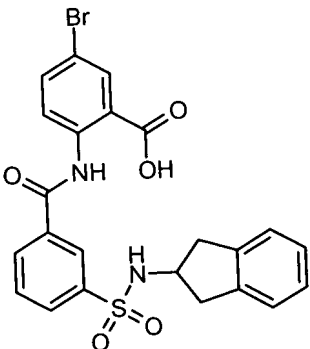
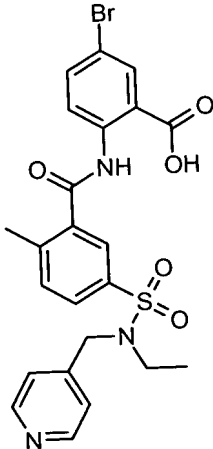
Compound No., Structure	Compound No., Structure
<p>PHA-500230</p> 	<p>PHA-500236</p> 
<p>PHA-500248</p> 	<p>PHA-500284</p> 
<p>PHA-502605</p> 	<p>PHA-502606</p> 
<p>PHA-520185</p> 	<p>PHA-520200</p> 

Compound No., Structure	Compound No., Structure
PHA-520221 	PHA-520245 
PHA-520412 	PHA-520413 
PHA-520414 	PHA-520416 
PHA-523506 	PHA-523507 

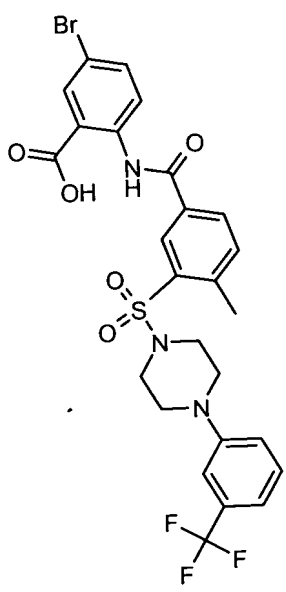
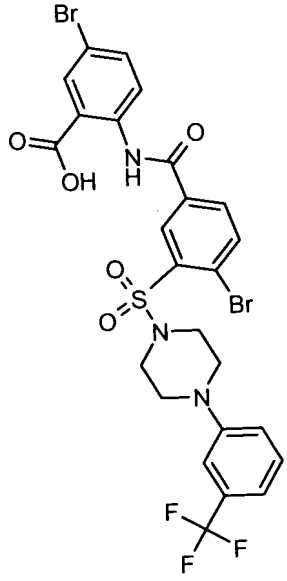
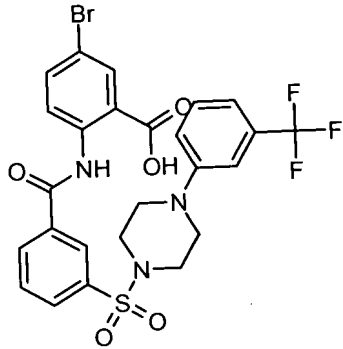
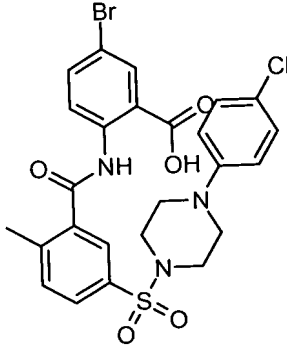
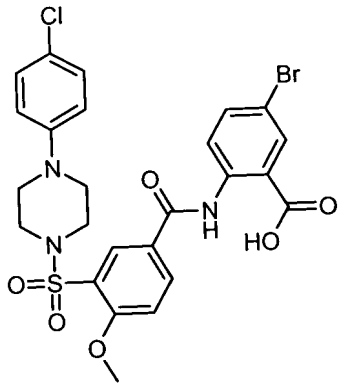
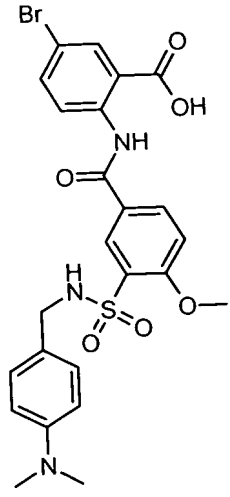
Compound No., Structure	Compound No., Structure
PHA-523508 	PHA-523509 
PHA-523510 	PHA-523511 
PHA-523512 	PHA-523513 
PHA-523514 	PHA-523515 
PHA-523516 	PHA-523517 

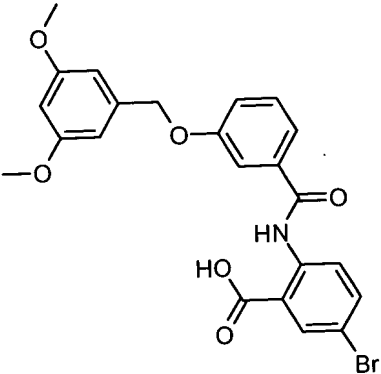
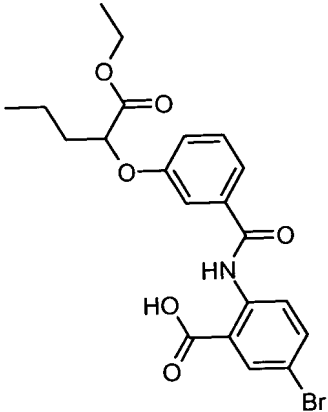
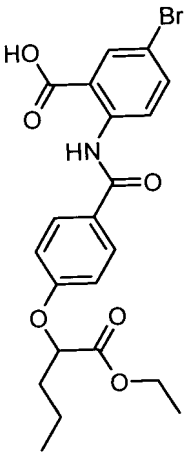
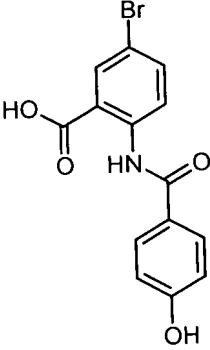
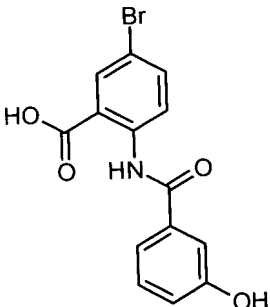
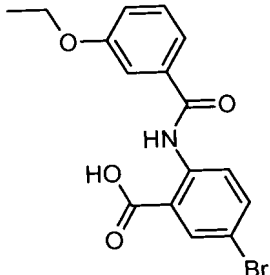
Compound No., Structure	Compound No., Structure
PHA-523518 	PHA-523519 
PHA-523520 	PHA-523521 
PHA-524545E 	PHA-524553A  <p>HCl</p>
PHA-525500 	PHA-525501 

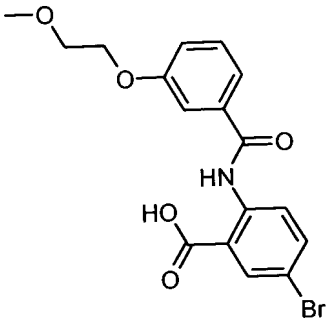
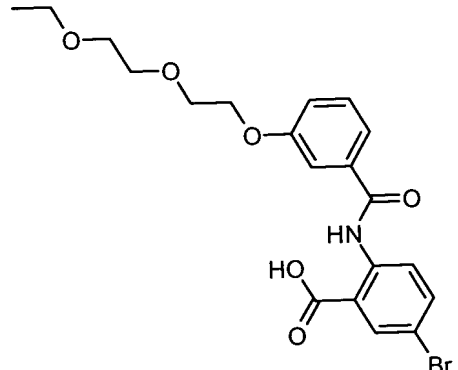
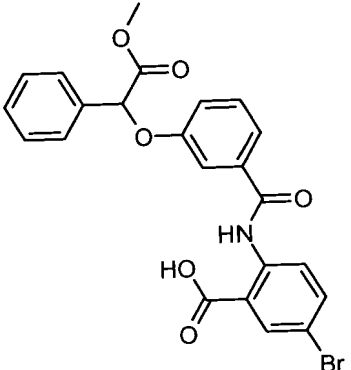
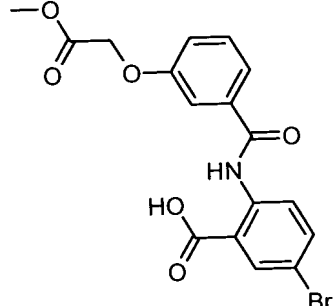
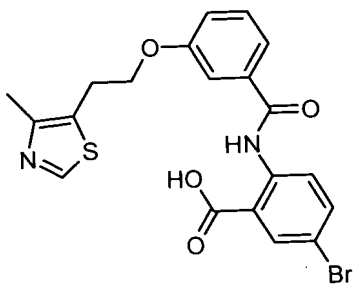
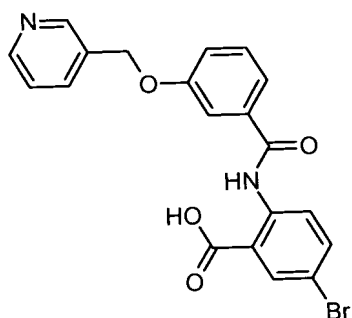
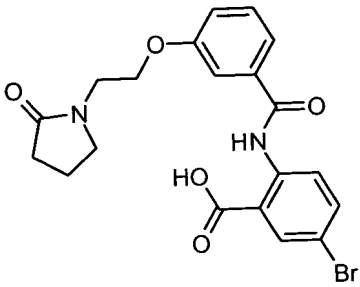
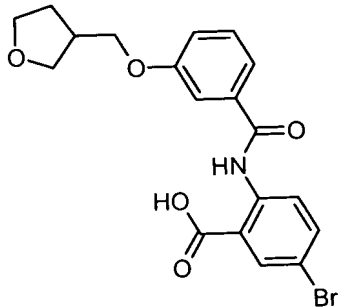
Compound No., Structure	Compound No., Structure
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<p data-bbox="235 609 414 651">PHA-525504</p> 	<p data-bbox="803 588 982 630">PHA-525505</p> 
<p data-bbox="251 1186 430 1228">PHA-525506</p> 	<p data-bbox="820 1165 998 1207">PHA-526641</p> 

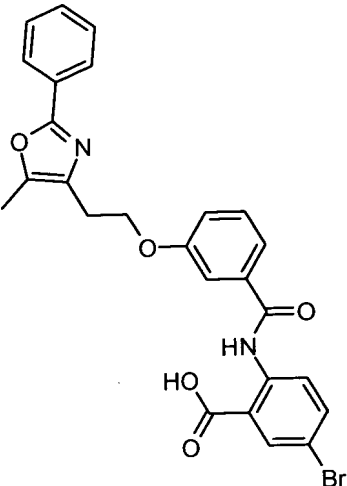
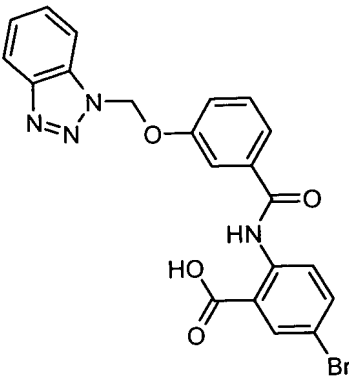
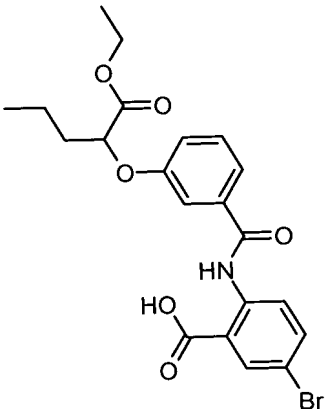
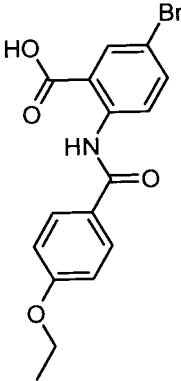
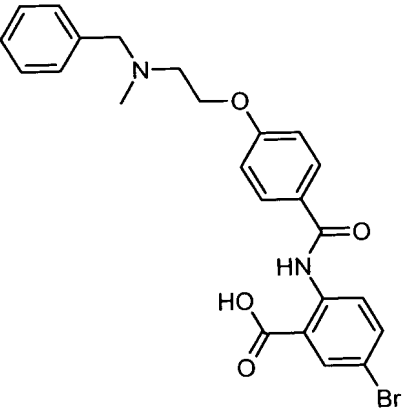
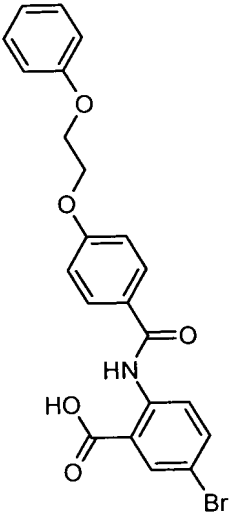
Compound No., Structure	Compound No., Structure
<p data-bbox="224 191 397 226">PHA-526643</p> 	<p data-bbox="792 170 966 205">PHA-526648</p> 
<p data-bbox="240 779 414 814">PHA-526650</p> 	<p data-bbox="808 758 982 793">PHA-526651</p> 
<p data-bbox="264 1367 438 1402">PHA-526652</p> 	<p data-bbox="833 1346 1006 1381">PHA-526653</p> 

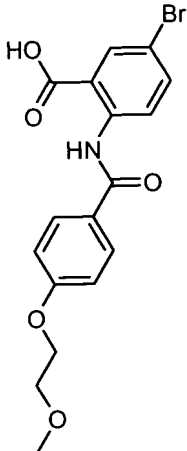
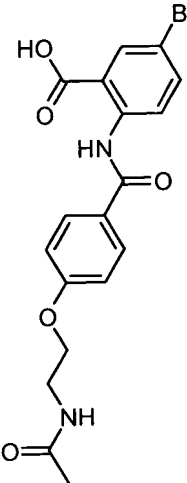
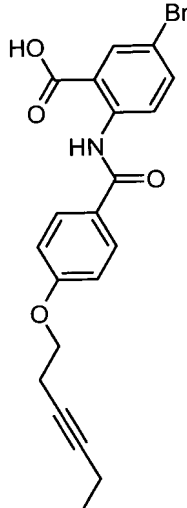
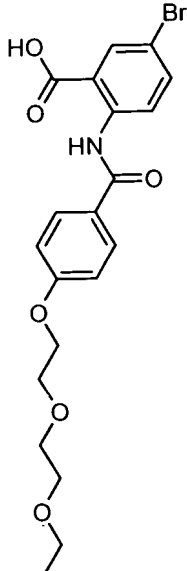
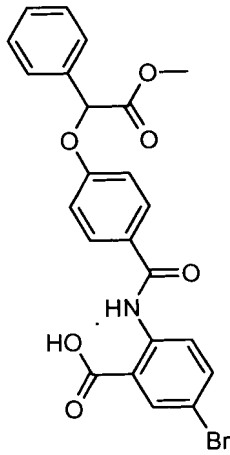
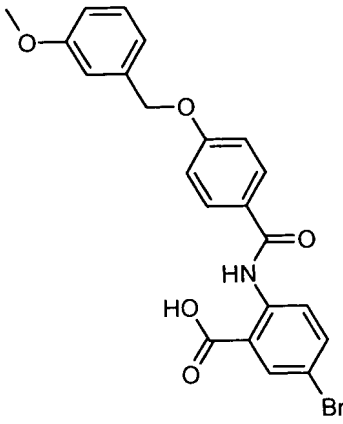


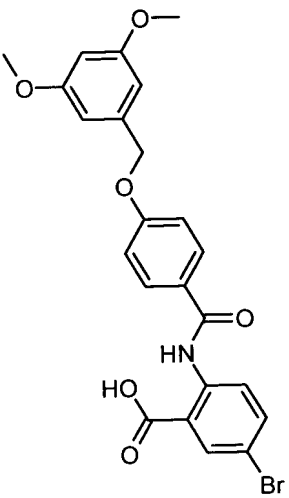
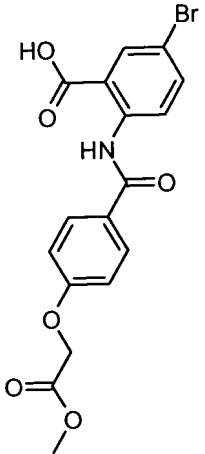
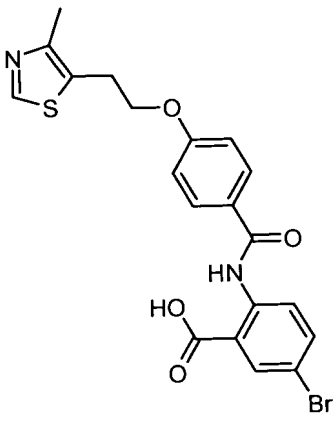
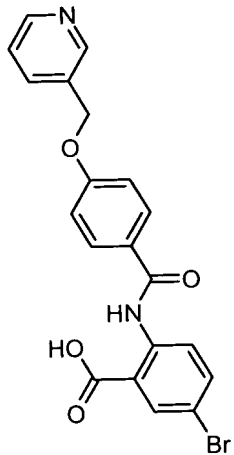
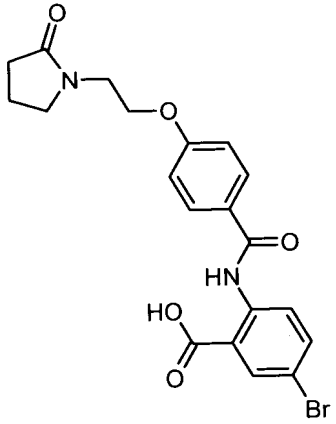
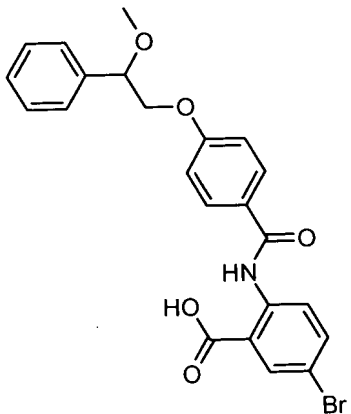
Compound No., Structure	Compound No., Structure
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<p data-bbox="243 871 417 905">PHA-526661</p> 	<p data-bbox="812 850 985 884">PHA-526679</p> 
<p data-bbox="259 1312 433 1346">PHA-526681</p> 	<p data-bbox="828 1291 1002 1325">PHA-526683</p> 

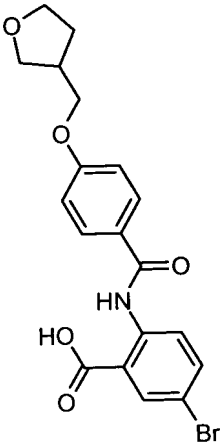
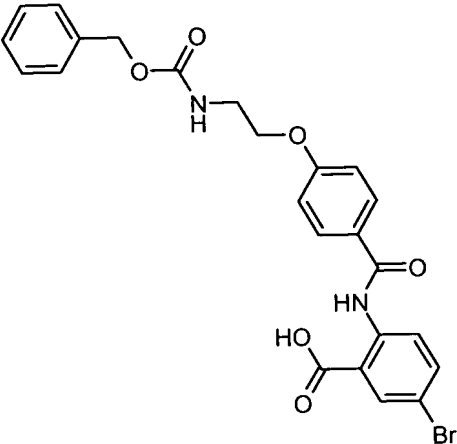
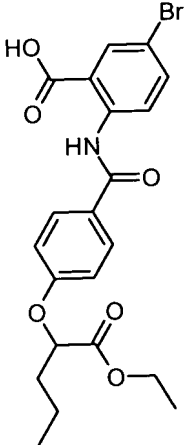
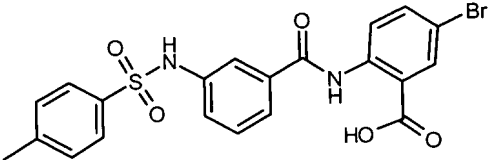
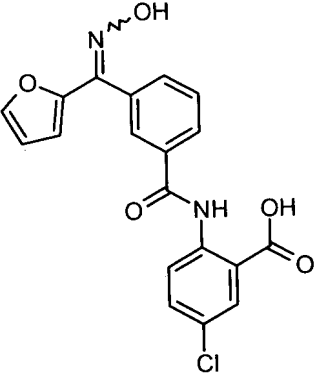
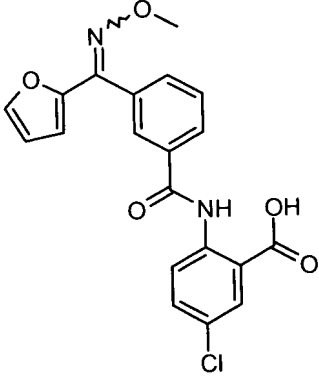
Compound No., Structure	Compound No., Structure
<p data-bbox="233 184 407 220">PHA-526705</p>  <chem data-bbox="329 254 708 625">COc1cc(OC)cc(COc2ccc(cc2)C(=O)Nc3ccc(cc3)C(=O)O)c1</chem>	<p data-bbox="800 170 974 205">PHA-526707</p>  <chem data-bbox="922 233 1247 642">CCCC(=O)Oc1ccc(cc1)C(=O)Nc2ccc(cc2)C(=O)OBr</chem>
<p data-bbox="248 709 422 745">PHA-526712</p>  <chem data-bbox="430 758 613 1209">CCCC(=O)Oc1ccc(cc1)C(=O)Nc2ccc(cc2)C(=O)OBr</chem>	<p data-bbox="816 688 990 724">PHA-530914</p>  <chem data-bbox="987 747 1195 1094">Oc1ccc(cc1)C(=O)Nc2ccc(cc2)C(=O)OBr</chem>
<p data-bbox="264 1255 438 1291">PHA-530915</p>  <chem data-bbox="410 1304 678 1608">Oc1ccc(cc1)C(=O)Nc2ccc(cc2)C(=O)OBr</chem>	<p data-bbox="833 1234 1006 1270">PHA-533232</p>  <chem data-bbox="979 1293 1247 1566">CCOc1ccc(cc1)C(=O)Nc2ccc(cc2)C(=O)OBr</chem>

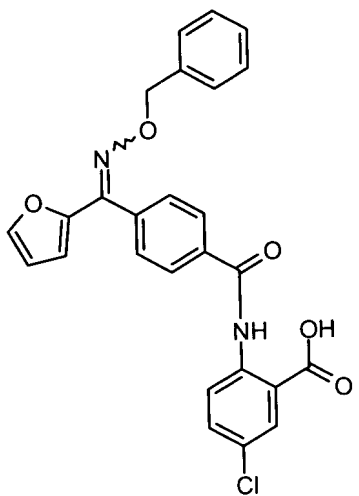
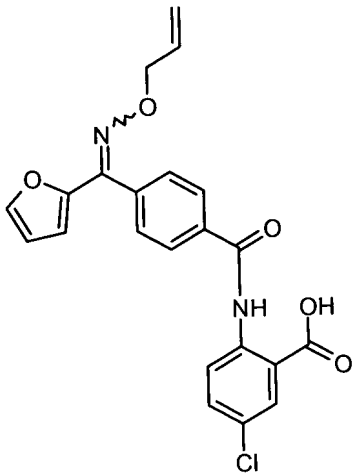
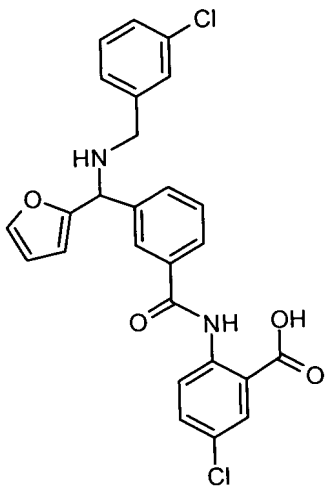
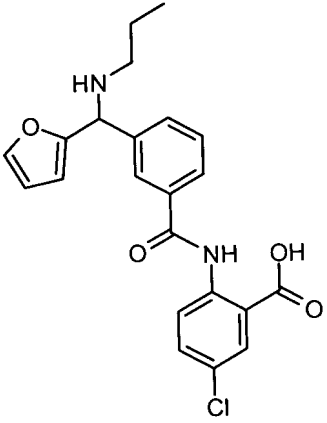
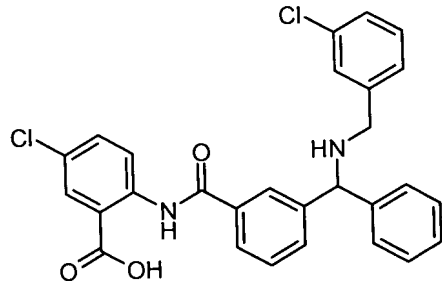
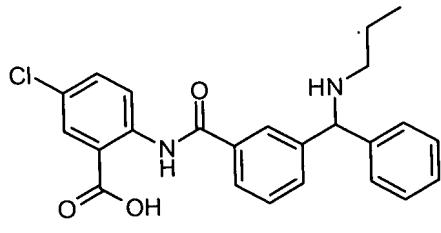
Compound No., Structure	Compound No., Structure
PHA-533237 	PHA-533243 
PHA-533244 	PHA-533247 
PHA-533249 	PHA-533252 
PHA-533253 	PHA-533257 

Compound No., Structure	Compound No., Structure
<p data-bbox="245 174 418 210">PHA-533258</p>  <chem data-bbox="354 241 699 724">Cc1cc(C2=CN(C2COc3ccc(cc3)C(=O)Nc4ccc(cc4)C(=O)O)cc5ccccc5)oc6ccccc6</chem>	<p data-bbox="813 168 987 203">PHA-533259</p>  <chem data-bbox="917 226 1263 604">BrC1=CC=C(C(=C1)C(=O)Nc2ccc(cc2)C(=O)O)C(=O)Nc3ccc(cc3)COc4nn[nH]4</chem>
<p data-bbox="256 768 430 804">PHA-533261</p>  <chem data-bbox="375 835 699 1241">CC(C)C(COC(=O)OCC)COc1ccc(cc1)C(=O)Nc2ccc(cc2)C(=O)O</chem>	<p data-bbox="824 762 998 798">PHA-533262</p>  <chem data-bbox="1003 814 1182 1192">BrC1=CC=C(C(=C1)C(=O)Nc2ccc(cc2)C(=O)O)C(=O)Nc3ccc(cc3)COCC</chem>
<p data-bbox="267 1287 441 1323">PHA-533264</p>  <chem data-bbox="349 1354 748 1759">BrC1=CC=C(C(=C1)C(=O)Nc2ccc(cc2)C(=O)O)C(=O)Nc3ccc(cc3)OCCN(C)Cc4ccccc4</chem>	<p data-bbox="836 1276 1010 1312">PHA-533265</p>  <chem data-bbox="1003 1339 1230 1850">BrC1=CC=C(C(=C1)C(=O)Nc2ccc(cc2)C(=O)O)C(=O)Nc3ccc(cc3)Oc4ccc(cc4)OCCOc5ccccc5</chem>

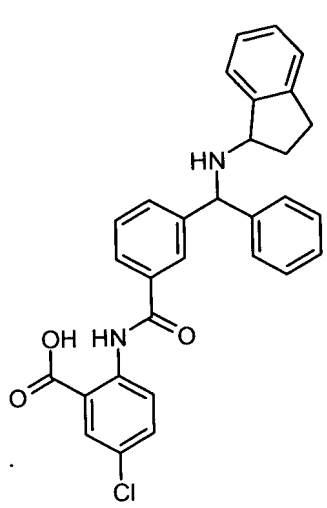
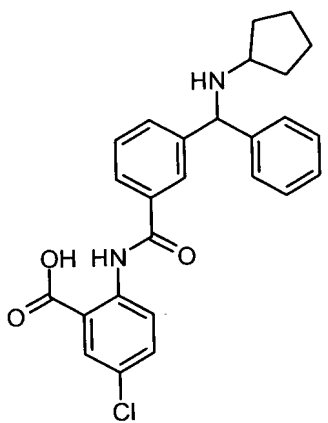
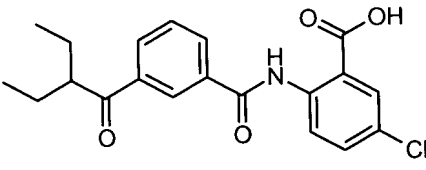
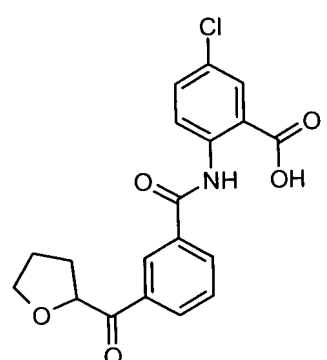
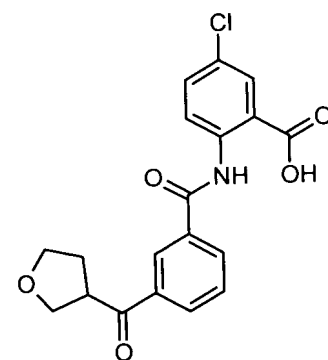
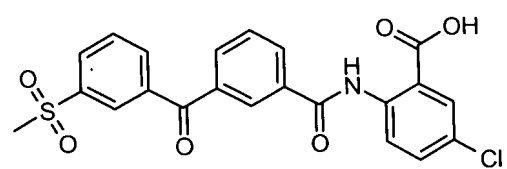
Compound No., Structure	Compound No., Structure
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<p data-bbox="256 758 431 789">PHA-533269</p> 	<p data-bbox="824 758 1000 789">PHA-533272</p> 
<p data-bbox="256 1430 431 1461">PHA-533273</p> 	<p data-bbox="824 1430 1000 1461">PHA-533274</p> 

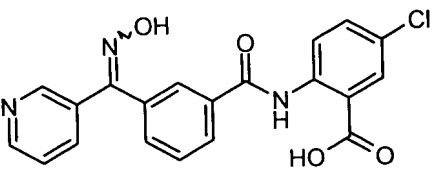
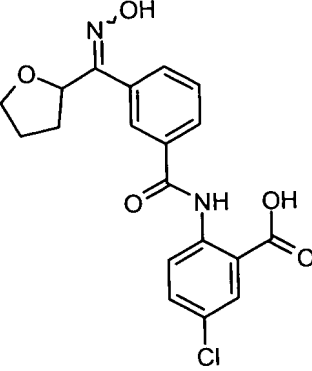
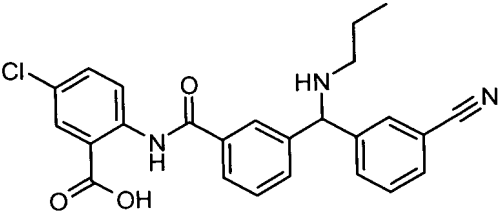
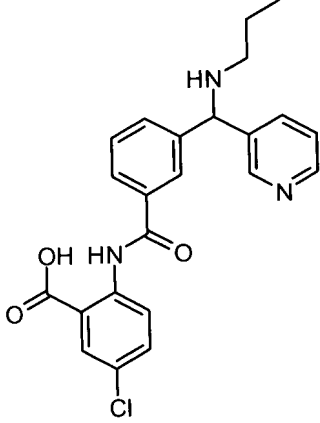
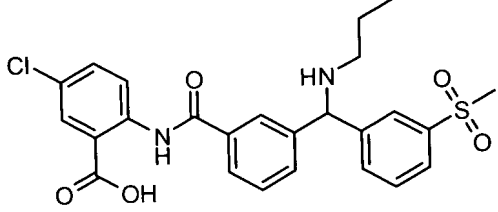
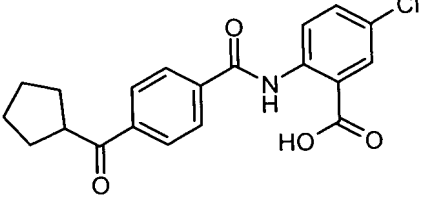
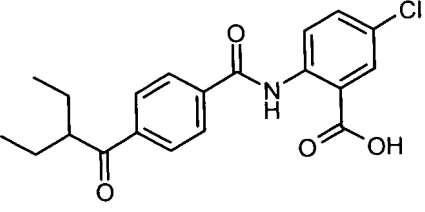
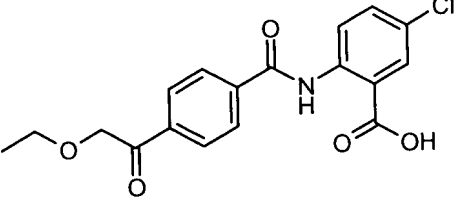
Compound No., Structure	Compound No., Structure
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<p>PHA-533278</p> 	<p>PHA-533281</p> 
<p>PHA-533282</p> 	<p>PHA-533285</p> 

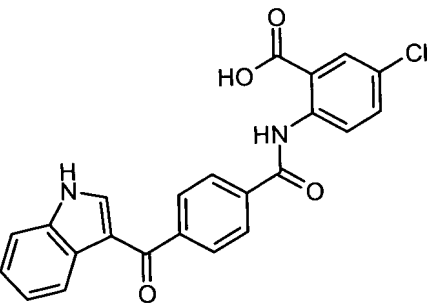
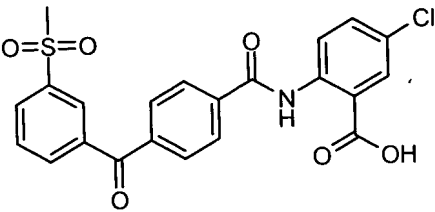
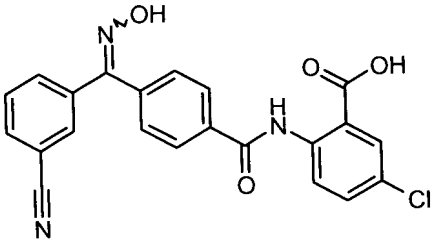
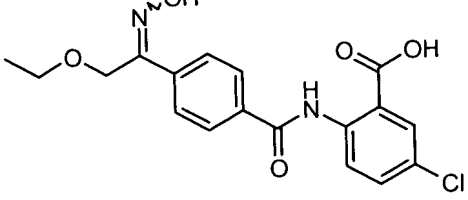
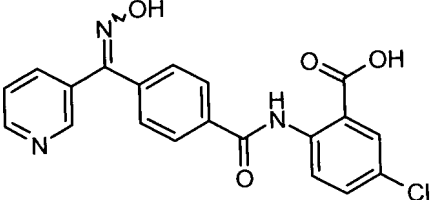
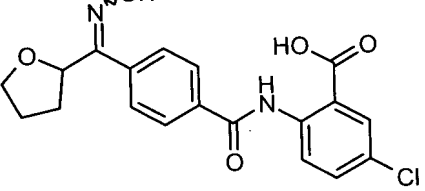
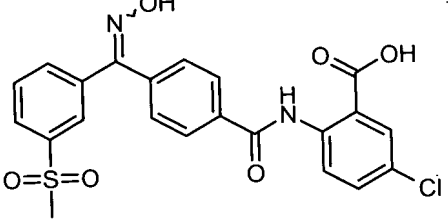
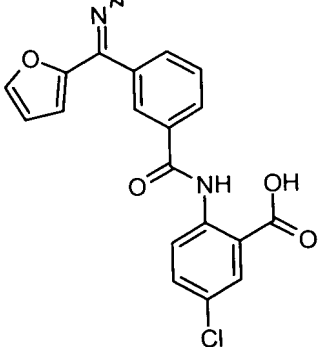
Compound No., Structure	Compound No., Structure
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<p data-bbox="245 730 431 762">PHA-533290</p> 	<p data-bbox="813 720 1000 751">PHA-533401</p> 
<p data-bbox="245 1276 448 1308">PHA-537084</p>  <p data-bbox="355 1719 761 1751">least retained isomer by RP-HPLC</p>	<p data-bbox="813 1266 1016 1297">PHA-537085</p> 

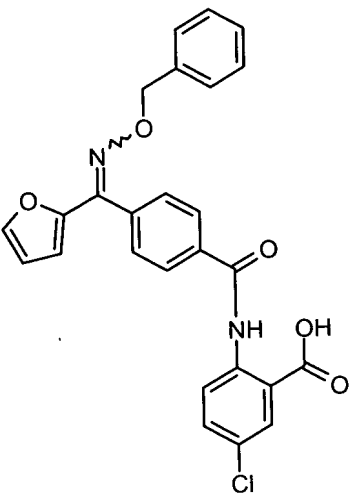
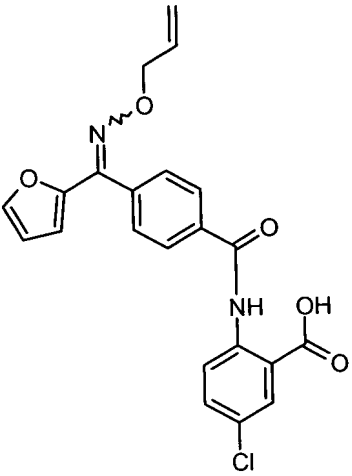
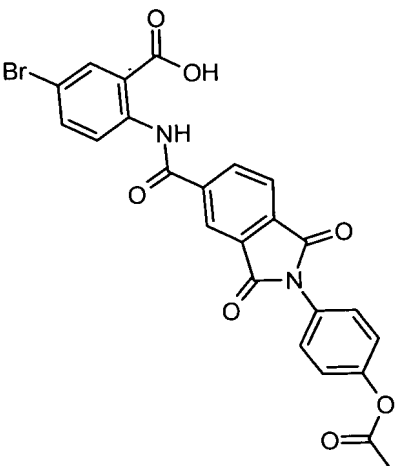
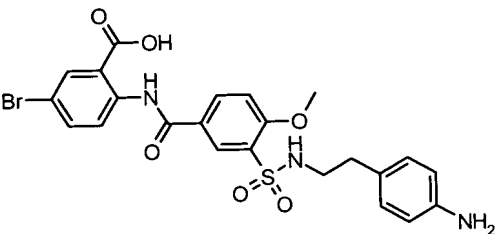
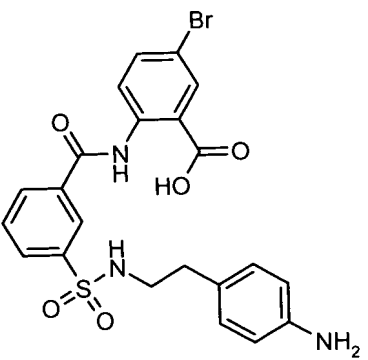
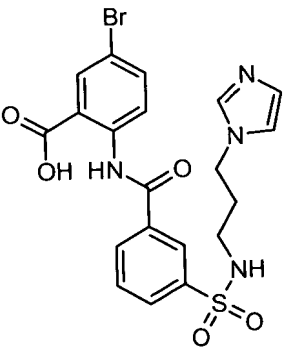
Compound No., Structure	Compound No., Structure
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<p data-bbox="251 798 430 829">PHA-537091</p> 	<p data-bbox="820 787 998 819">PHA-537092</p> 
<p data-bbox="267 1375 446 1407">PHA-537098</p> 	<p data-bbox="836 1365 1015 1396">PHA-537099</p> 

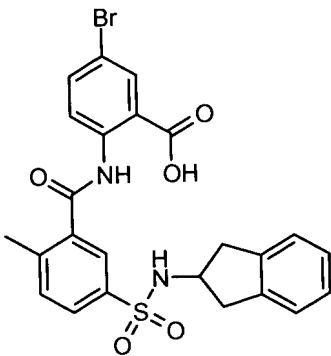
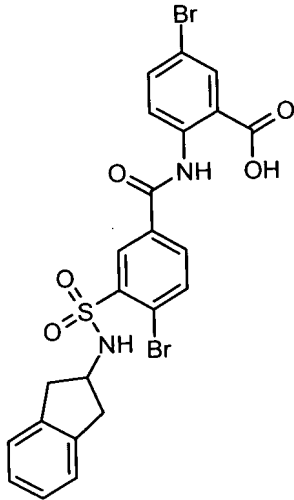
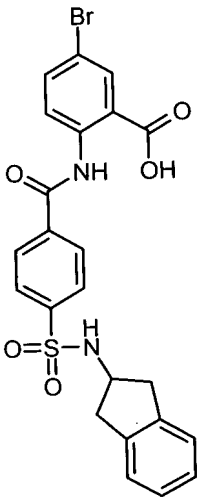
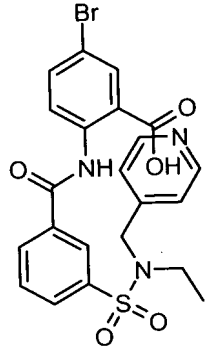
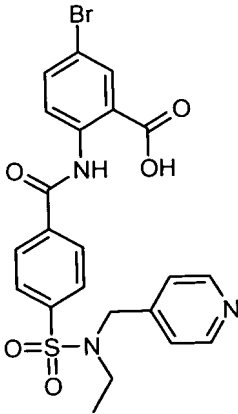
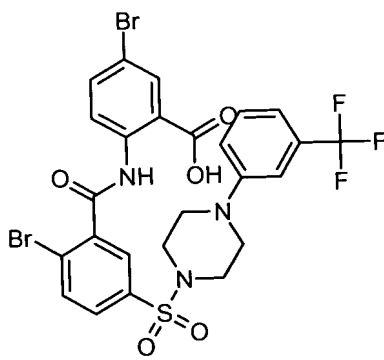


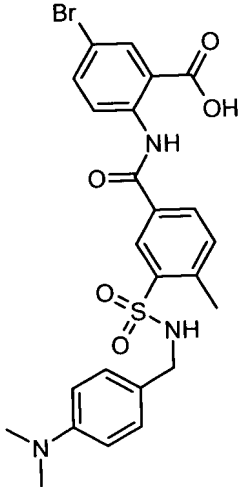
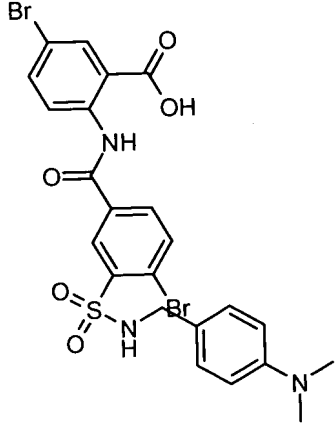
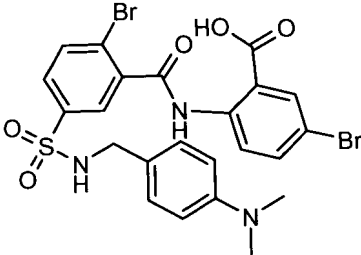
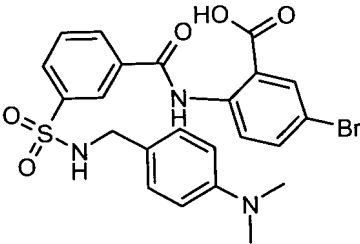
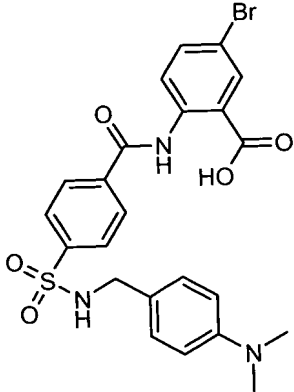
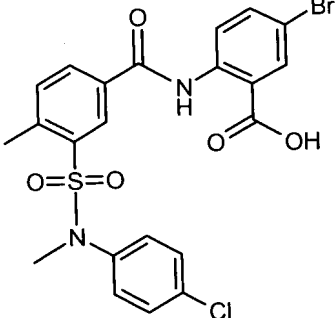
Compound No., Structure	Compound No., Structure
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<p>PHA-537106</p> 	<p>PHA-537110</p> 
<p>PHA-537112</p> 	<p>PHA-537114</p> 

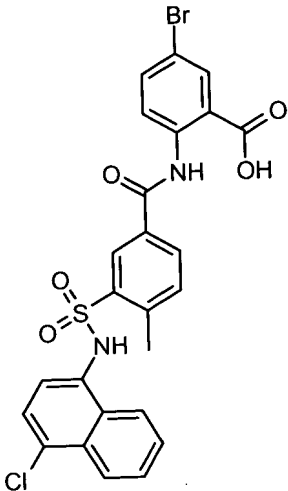
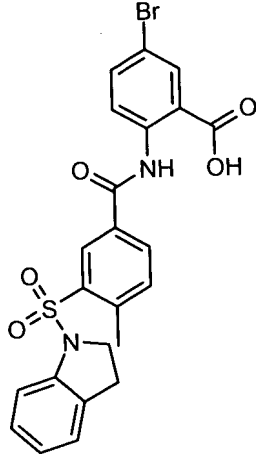
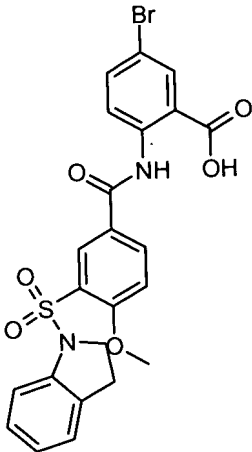
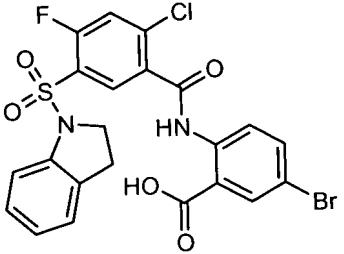
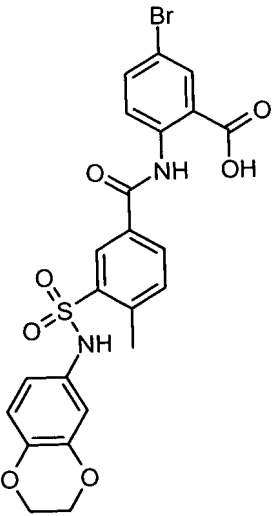
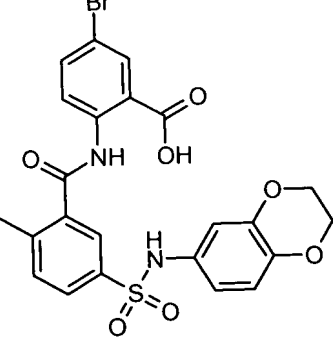
Compound No., Structure	Compound No., Structure
<p>PHA-537121</p> 	<p>PHA-537122</p> 
<p>PHA-537128</p> 	<p>PHA-537133</p> 
<p>PHA-537138</p> 	<p>PHA-537139</p> 
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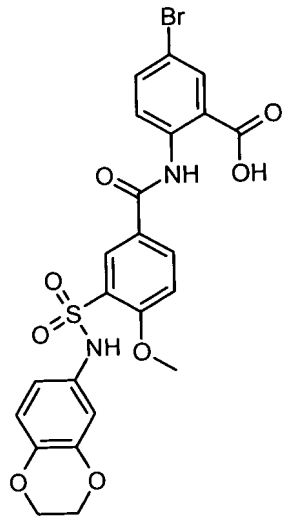
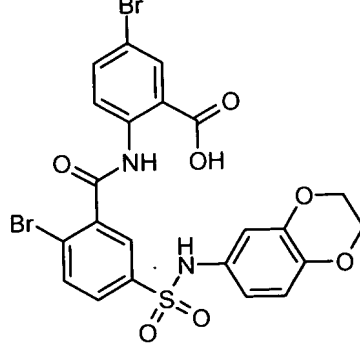
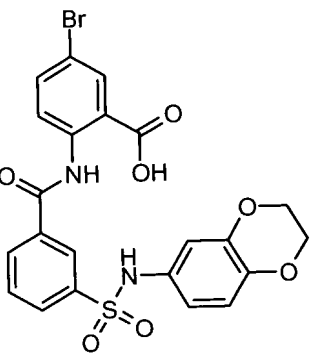
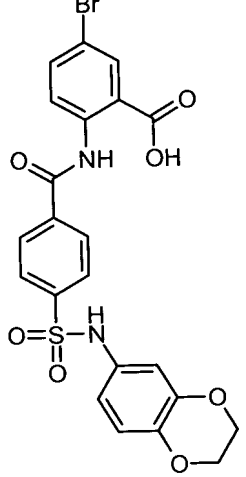
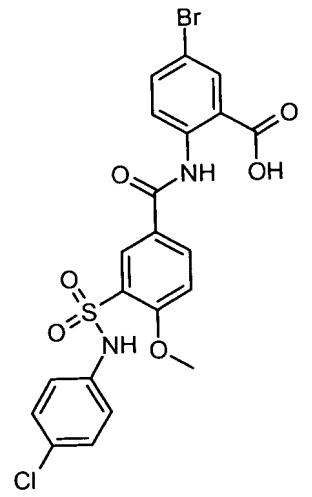
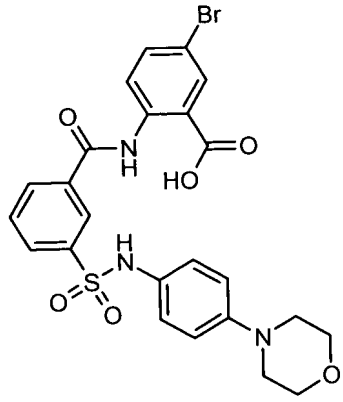
Compound No., Structure	Compound No., Structure
PHA-537144 	PHA-537150 
PHA-537152 	PHA-537155 
PHA-537157 	PHA-537158 
PHA-537162 	PHA-537202  most highly retained isomer by RP-LC/MS

Compound No., Structure	Compound No., Structure
<p>PHA-537203</p>  <p>most highly retained isomer by RP-LC/MS</p>	<p>PHA-537204</p>  <p>most highly retained isomer by RP-LC/M</p>
<p>PHA-538016</p> 	<p>PHA-539146</p> 
<p>PHA-539148</p> 	<p>PHA-539149</p> 

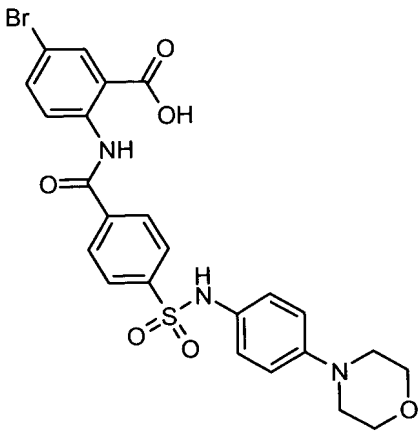
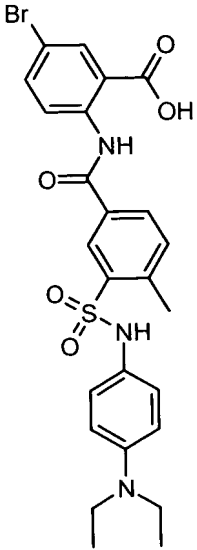
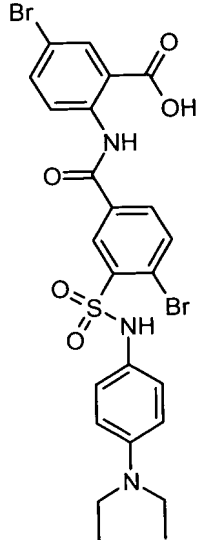
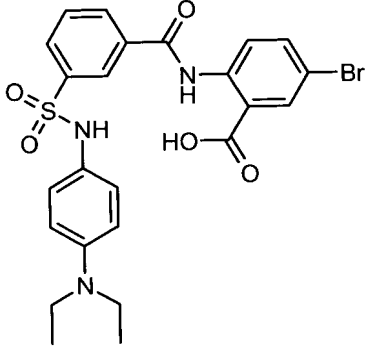
Compound No., Structure	Compound No., Structure
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<p data-bbox="256 768 430 800">PHA-539153</p> 	<p data-bbox="824 768 998 800">PHA-539154</p> 
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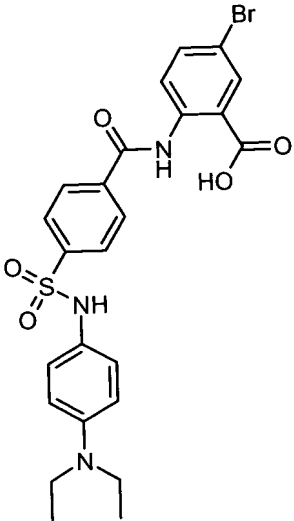
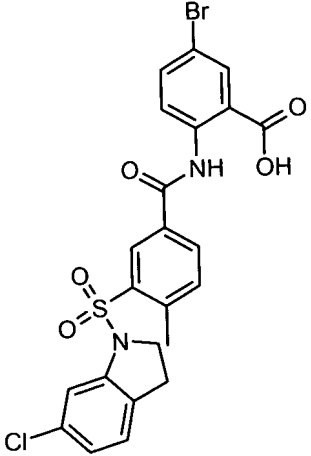
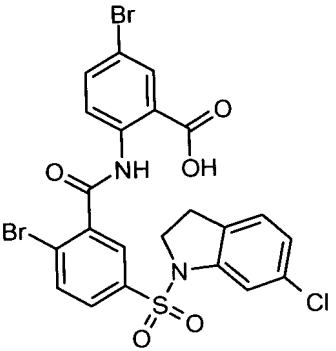
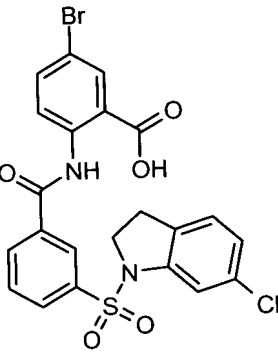
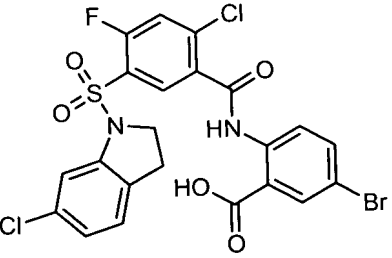
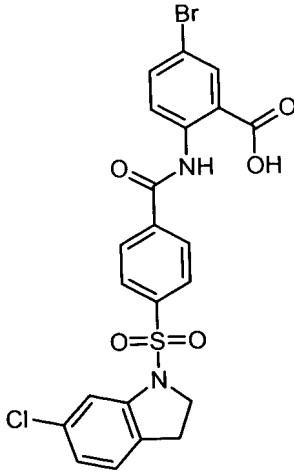
Compound No., Structure	Compound No., Structure
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<p data-bbox="256 762 431 800">PHA-539169</p> 	<p data-bbox="824 751 1000 789">PHA-539170</p> 
<p data-bbox="264 1119 440 1157">PHA-539171</p> 	<p data-bbox="833 1108 1008 1146">PHA-539172</p> 

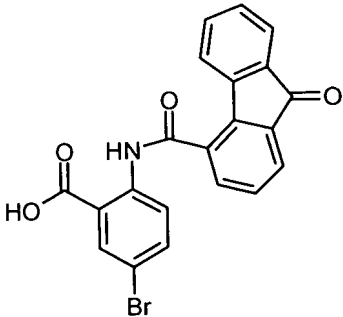
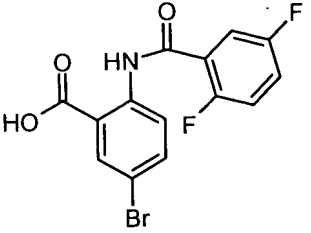
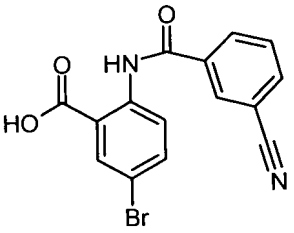
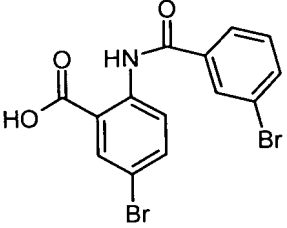
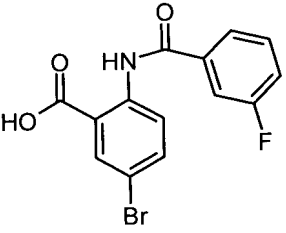
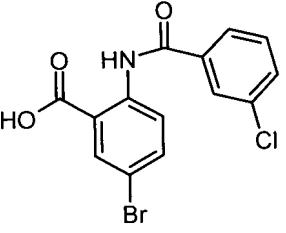
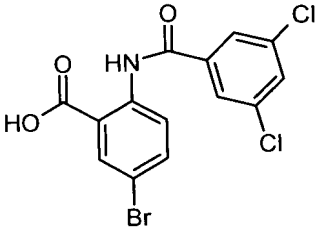
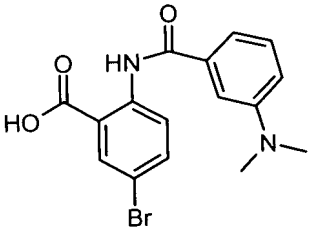
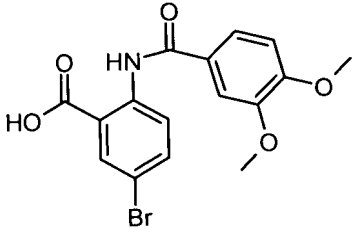
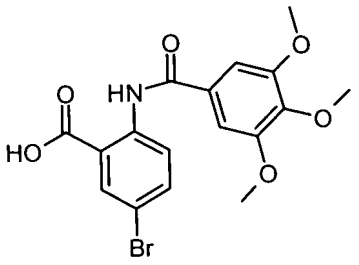
Compound No., Structure	Compound No., Structure
<p data-bbox="240 176 412 218">PHA-539174</p> 	<p data-bbox="808 176 980 218">PHA-539175</p> 
<p data-bbox="240 743 428 785">PHA-539177</p> 	<p data-bbox="808 743 997 785">PHA-539179</p> 
<p data-bbox="240 1310 444 1352">PHA-539180</p> 	<p data-bbox="808 1310 1013 1352">PHA-539181</p> 

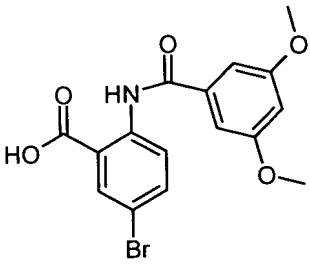
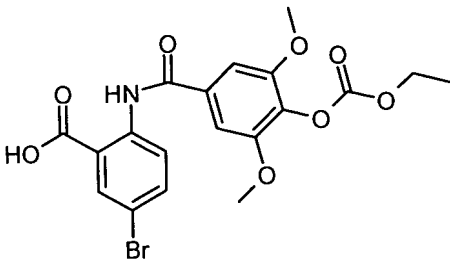
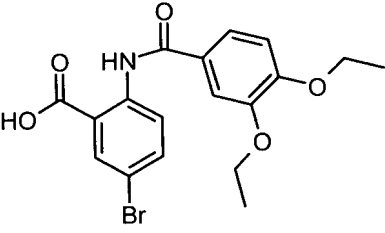
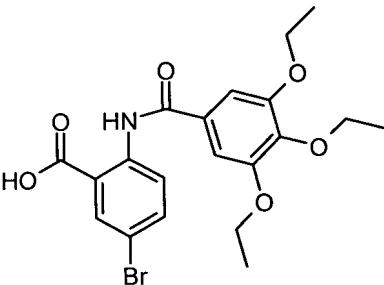
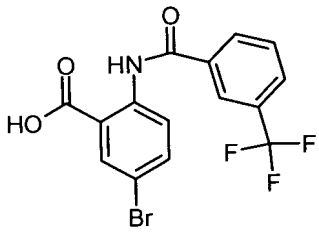
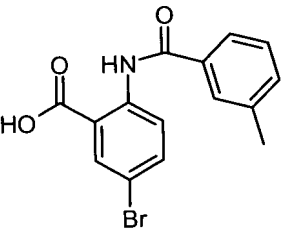
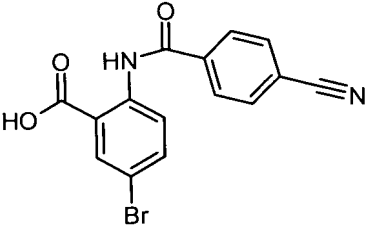
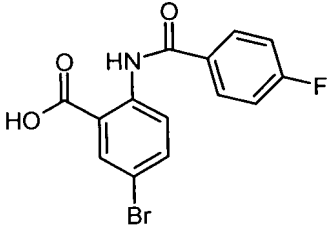
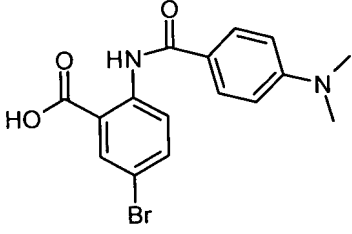
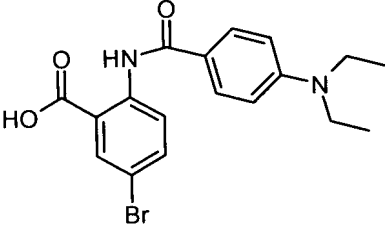
Compound No., Structure	Compound No., Structure
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<p data-bbox="251 777 430 819">PHA-539187</p> 	<p data-bbox="820 777 998 819">PHA-539188</p> 
<p data-bbox="251 1354 430 1396">PHA-539190</p> 	<p data-bbox="820 1354 998 1396">PHA-539193</p> 

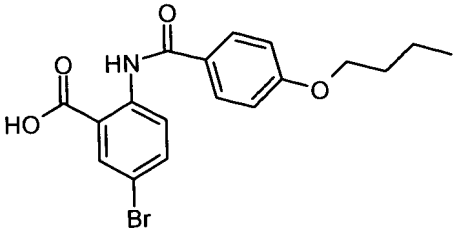
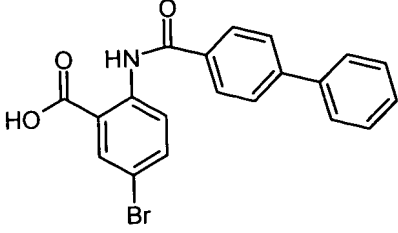
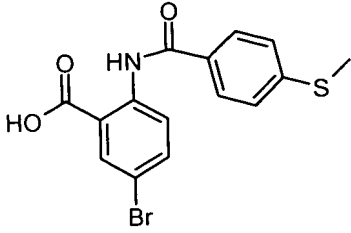
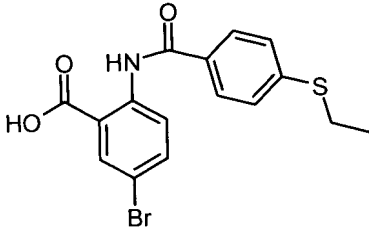
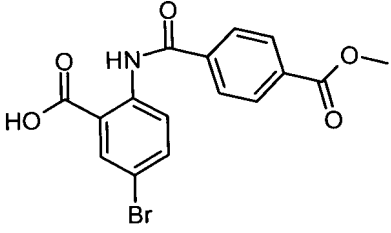
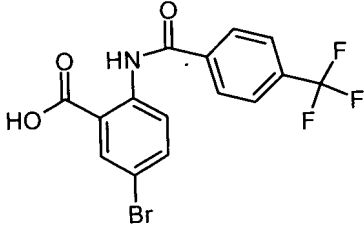
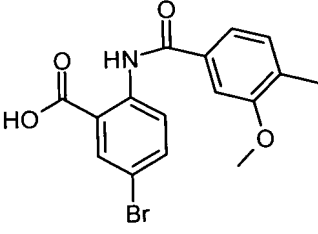
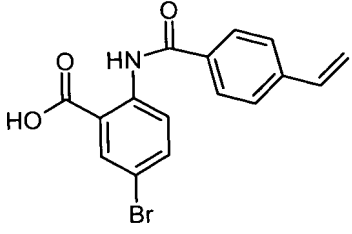
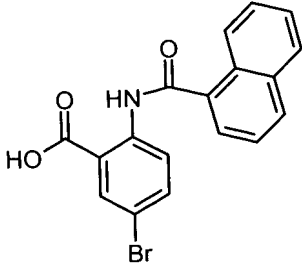
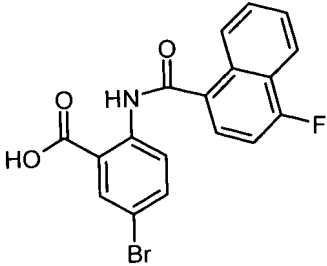


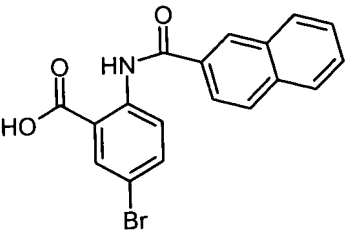
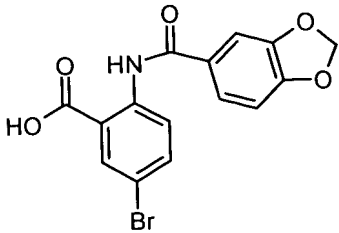
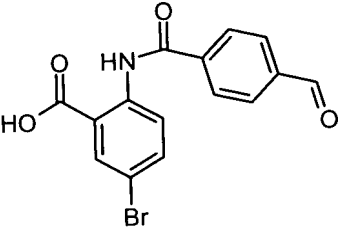
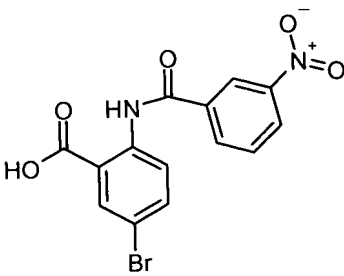
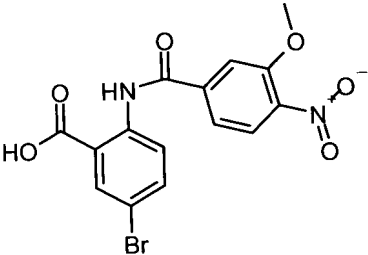
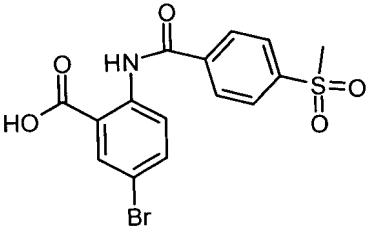
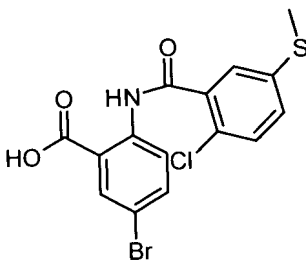
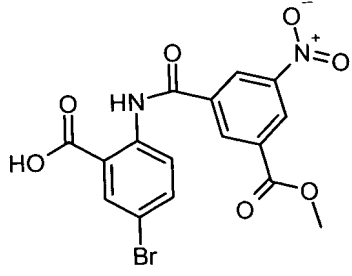
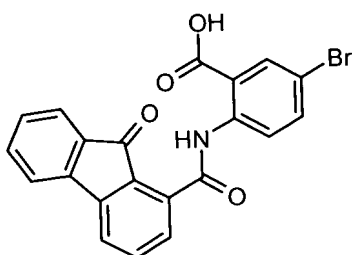
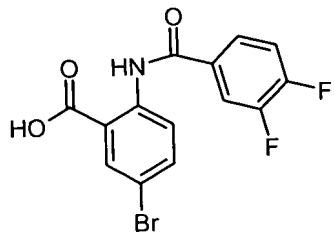
Compound No., Structure	Compound No., Structure
<p data-bbox="256 163 430 195">PHA-539194</p> 	<p data-bbox="824 153 998 184">PHA-539195</p> 
<p data-bbox="264 814 438 846">PHA-539197</p> 	<p data-bbox="833 804 1006 835">PHA-539198</p> 

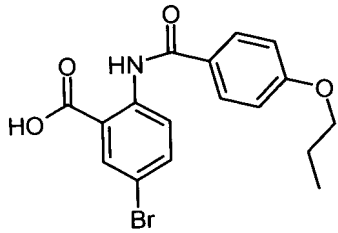
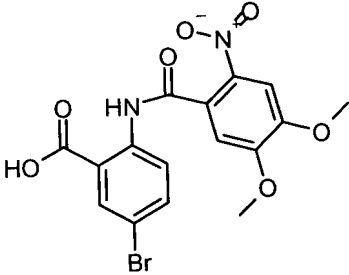
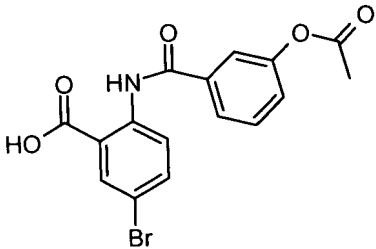
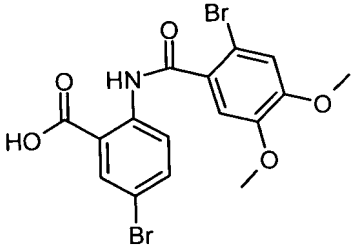
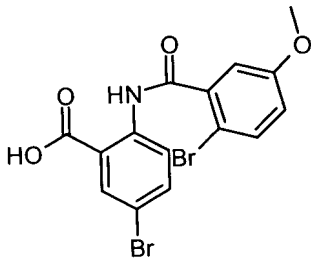
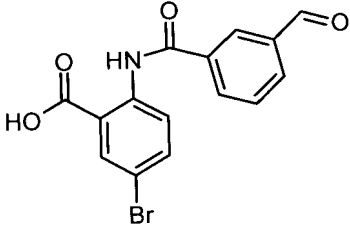
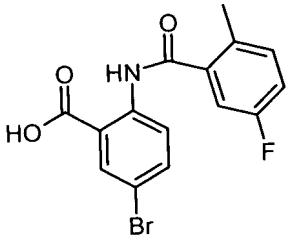
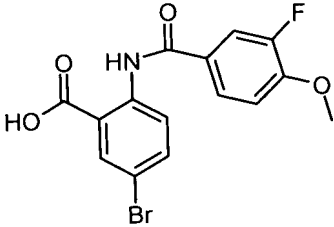
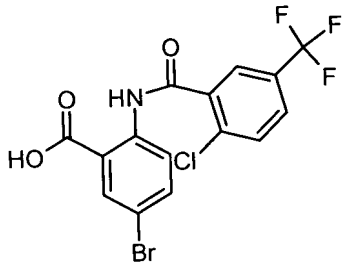
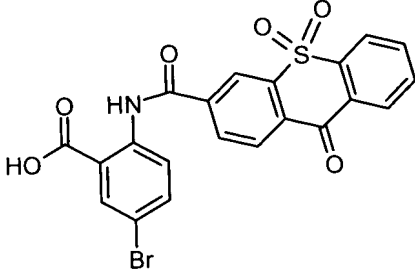
Compound No., Structure	Compound No., Structure
<p data-bbox="240 176 415 210">PHA-539199</p> 	<p data-bbox="808 155 984 189">PHA-539203</p> 
<p data-bbox="256 785 431 819">PHA-539206</p> 	<p data-bbox="824 764 1000 798">PHA-539207</p> 
<p data-bbox="272 1226 448 1260">PHA-539208</p> 	<p data-bbox="841 1205 1016 1239">PHA-539209</p> 

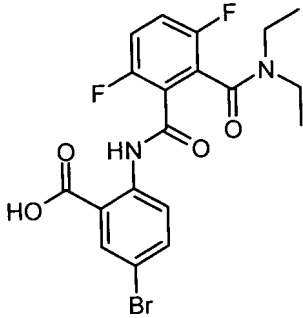
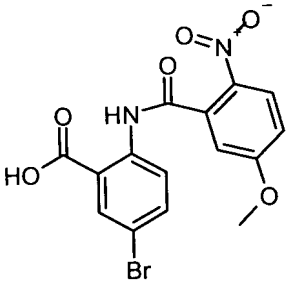
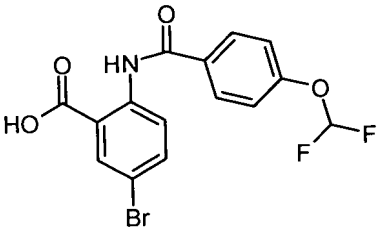
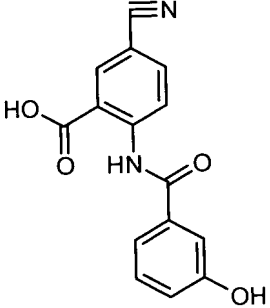
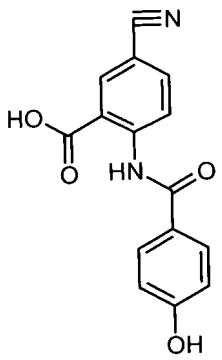
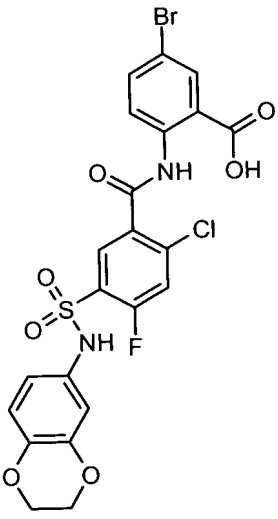
Compound No., Structure	Compound No., Structure
<p>PHA-539234</p> 	<p>PHA-539235</p> 
<p>PHA-539245</p> 	<p>PHA-539246</p> 
<p>PHA-539247</p> 	<p>PHA-539248</p> 
<p>PHA-539249</p> 	<p>PHA-539250</p> 
<p>PHA-539251</p> 	<p>PHA-539252</p> 

Compound No., Structure	Compound No., Structure
PHA-539253 	PHA-539254 
PHA-539255 	PHA-539256 
PHA-539257 	PHA-539258 
PHA-539259 	PHA-539260 
PHA-539262 	PHA-539263 

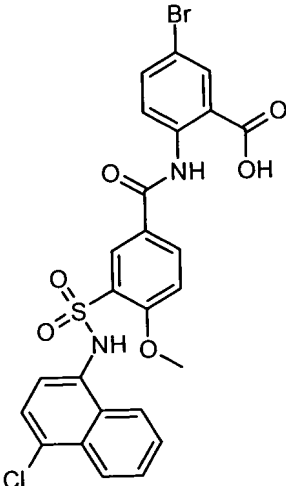
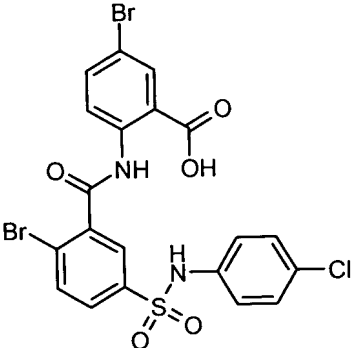
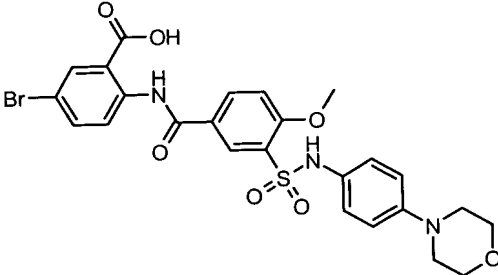
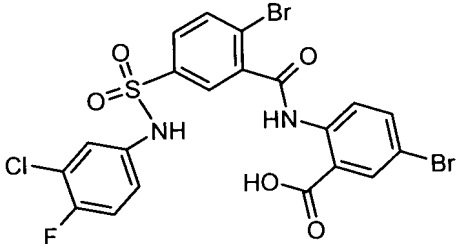
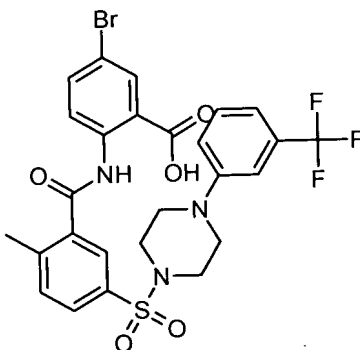
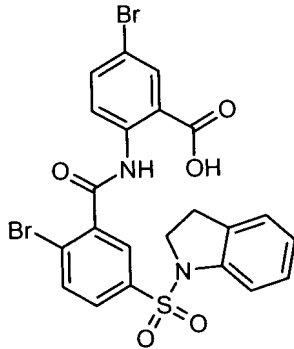
Compound No., Structure	Compound No., Structure
<p>PHA-539264</p> 	<p>PHA-539265</p> 
<p>PHA-539266</p> 	<p>PHA-539267</p> 
<p>PHA-539268</p> 	<p>PHA-539269</p> 
<p>PHA-539270</p> 	<p>PHA-539271</p> 
<p>PHA-539276</p> 	<p>PHA-539277</p> 

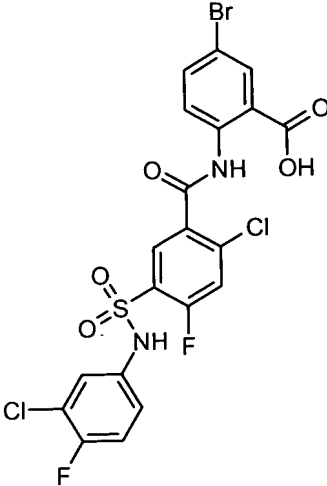
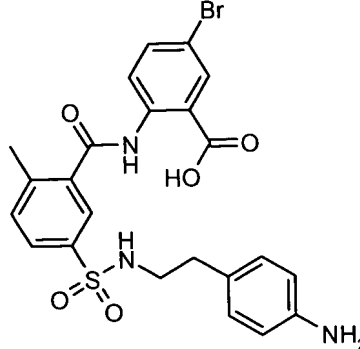
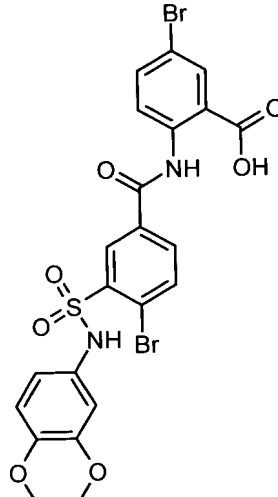
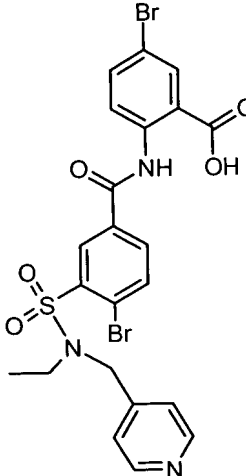
Compound No., Structure	Compound No., Structure
PHA-539278 	PHA-539285 
PHA-539293 	PHA-539294 
PHA-539295 	PHA-539296 
PHA-539297 	PHA-539298 
PHA-539302 	PHA-539303 

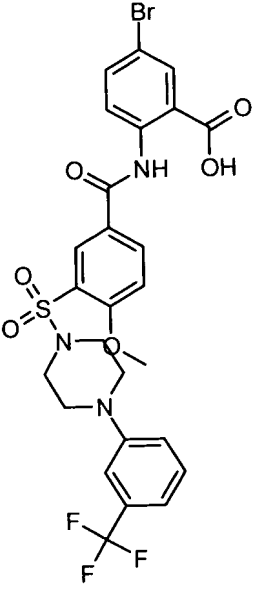
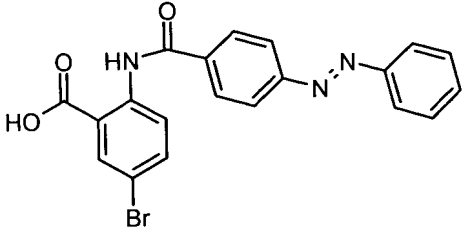
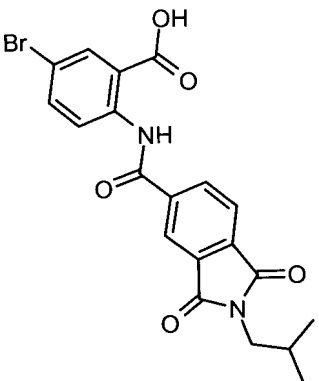
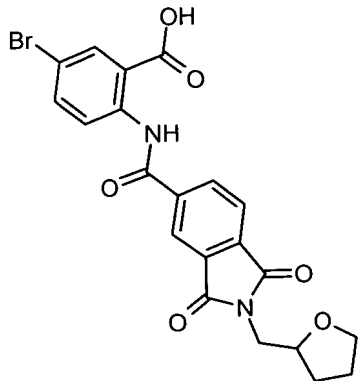
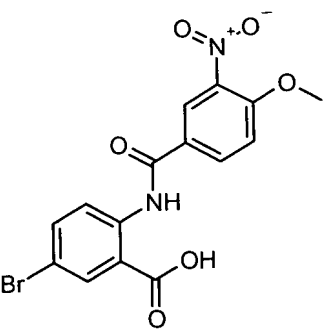
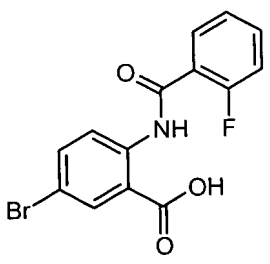
Compound No., Structure	Compound No., Structure
PHA-539305 	PHA-539307 
PHA-539308 	PHA-539310 
PHA-539312 	PHA-539313 
PHA-539314 	PHA-539317 
PHA-539318 	PHA-539322 

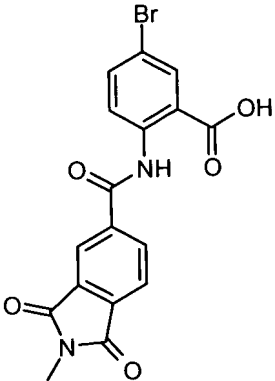
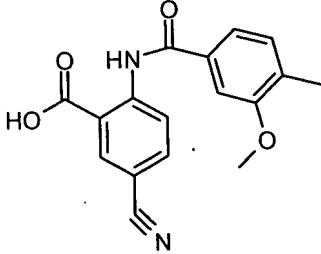
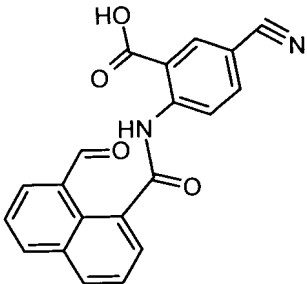
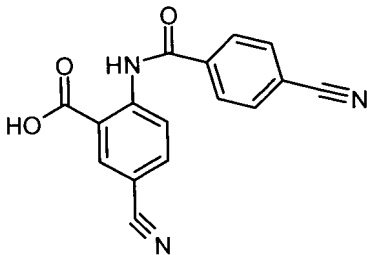
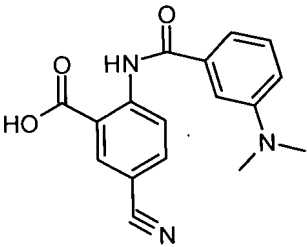
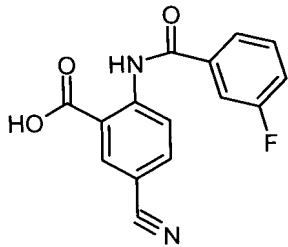
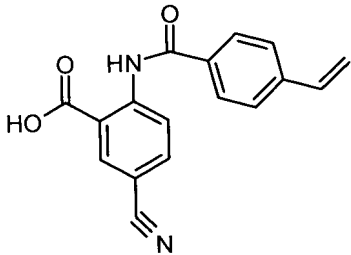
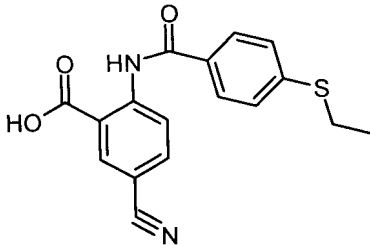
Compound No., Structure	Compound No., Structure
<p>PHA-539328</p> 	<p>PHA-539329</p> 
<p>PHA-539332</p> 	<p>PHA-539337</p> 
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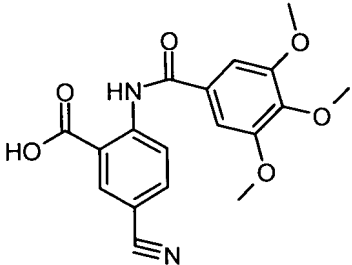
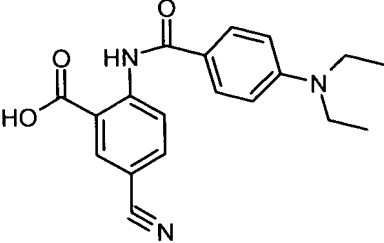
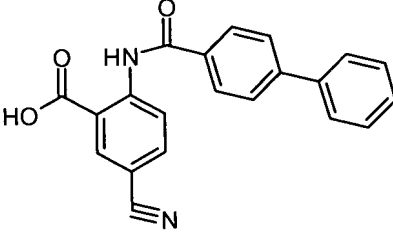
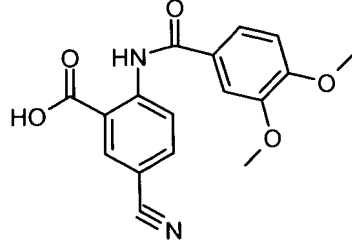
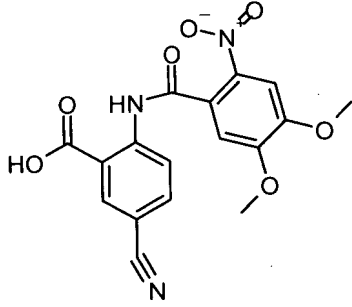
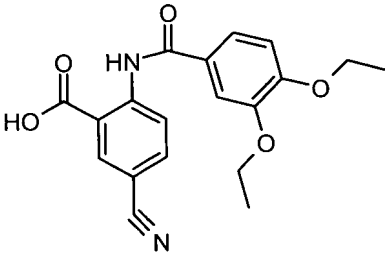
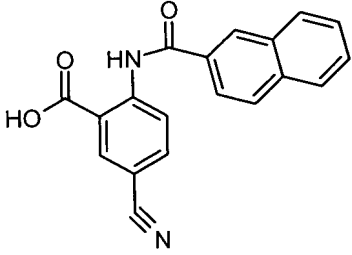
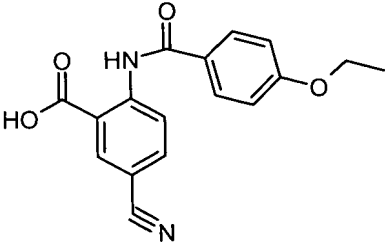


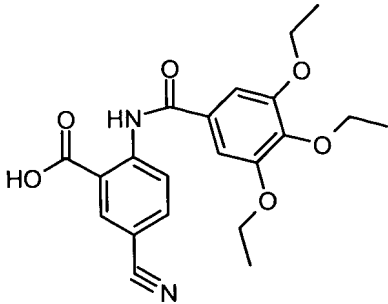
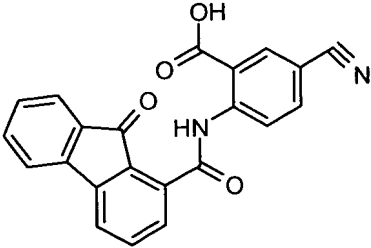
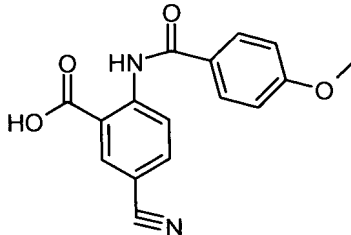
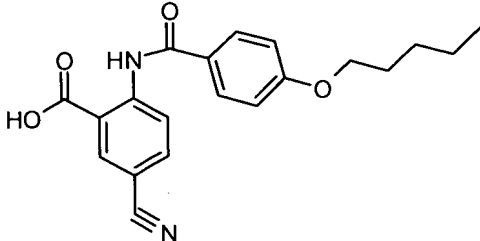
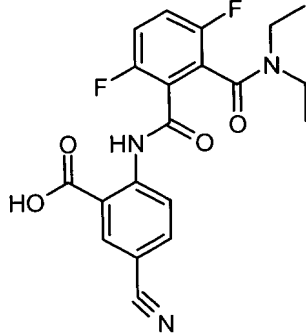
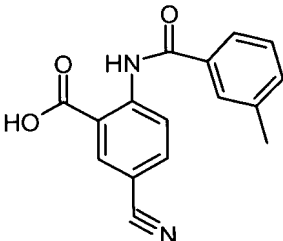
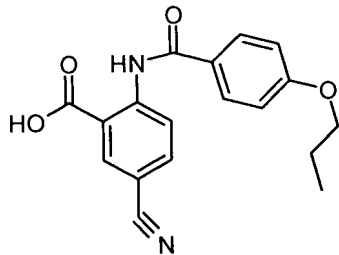
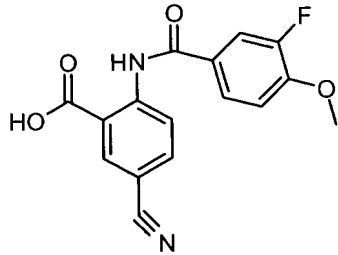
Compound No., Structure	Compound No., Structure
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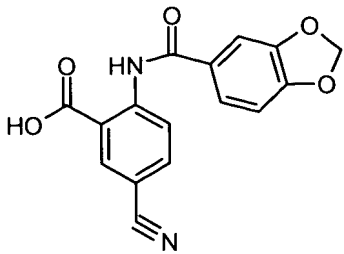
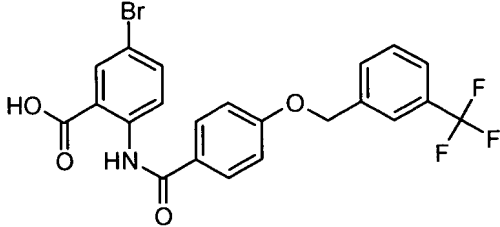
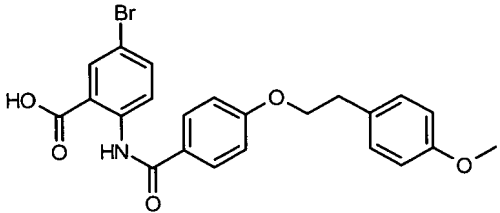
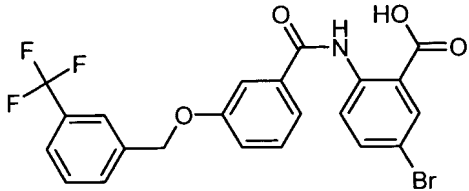
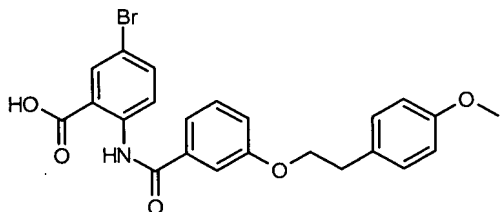
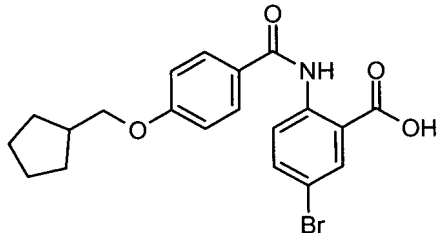
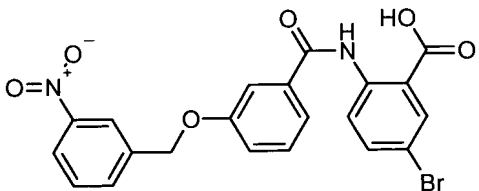
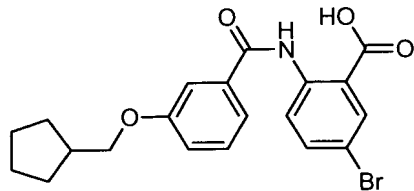
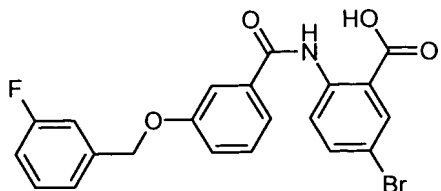
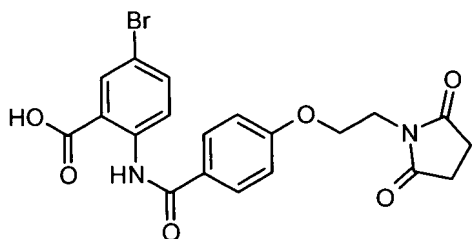
Compound No., Structure	Compound No., Structure
<p data-bbox="256 163 430 195">PHA-543695</p>  <p>The structure of PHA-543695 consists of a central benzene ring substituted with a carboxamide group (NH-C(=O)-) at position 1, a chlorine atom at position 2, a fluorine atom at position 3, and a sulfonamide group (-SO<sub>2</sub>NH-) at position 4. The sulfonamide group is further substituted with a 3-chloro-4-fluorophenyl ring.</p>	<p data-bbox="824 153 998 184">PHA-543698</p>  <p>The structure of PHA-543698 features a central benzene ring with a carboxamide group (NH-C(=O)-) at position 1, a hydroxyl group (-OH) at position 2, and a sulfonamide group (-SO<sub>2</sub>NH-) at position 3. The sulfonamide group is substituted with a 4-aminophenyl ring. Additionally, there is a methyl group at position 4 and a 4-bromophenyl ring at position 5 of the central benzene ring.</p>
<p data-bbox="264 741 438 772">PHA-543700</p>  <p>The structure of PHA-543700 is a benzene ring substituted with a carboxamide group (NH-C(=O)-) at position 1, a bromine atom at position 2, and a sulfonamide group (-SO<sub>2</sub>NH-) at position 3. The sulfonamide group is substituted with a 4-bromophenyl ring. The sulfonamide nitrogen is also part of a 1,3-benzodioxole system.</p>	<p data-bbox="833 730 1006 762">PHA-543701</p>  <p>The structure of PHA-543701 is a benzene ring substituted with a carboxamide group (NH-C(=O)-) at position 1, a bromine atom at position 2, and a sulfonamide group (-SO<sub>2</sub>NH-) at position 3. The sulfonamide group is substituted with a 4-pyridyl ring. The sulfonamide nitrogen is also substituted with an ethyl group.</p>

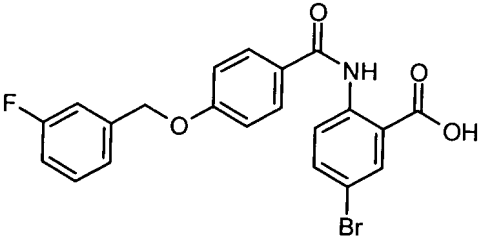
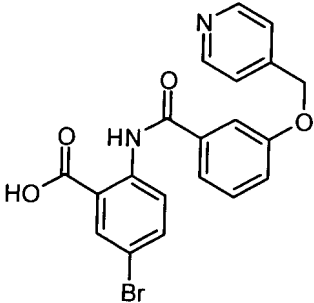
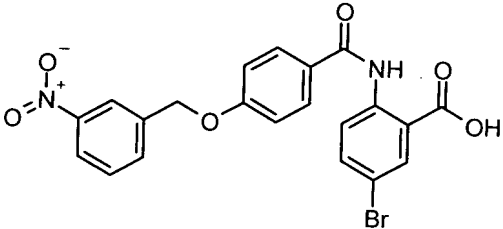
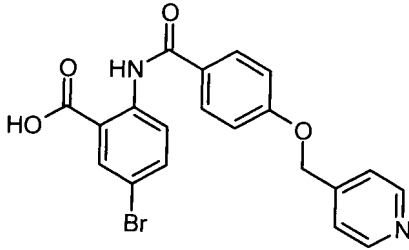
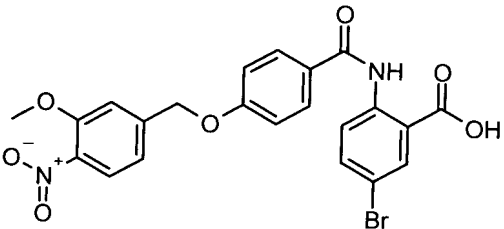
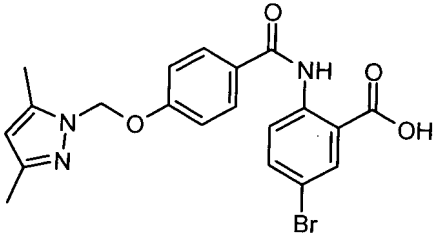
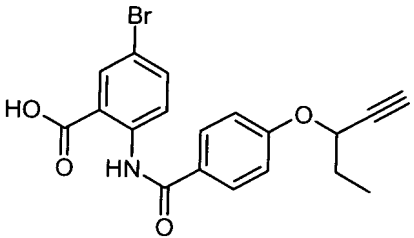
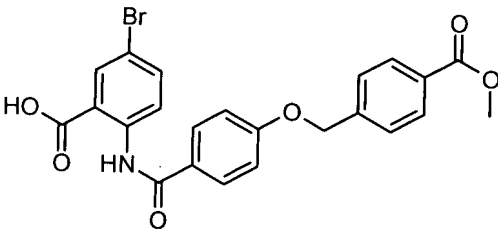
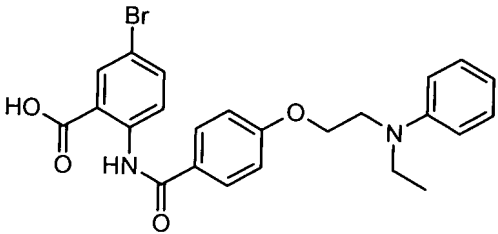
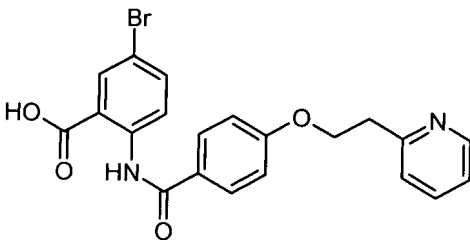
Compound No., Structure	Compound No., Structure
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<p>PHA-551675</p> 	<p>PHA-551716</p> 

Compound No., Structure	Compound No., Structure
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<p>PHA-563331</p> 	<p>PHA-563333</p> 
<p>PHA-563335</p> 	<p>PHA-563340</p> 
<p>PHA-563341</p> 	<p>PHA-563342</p> 

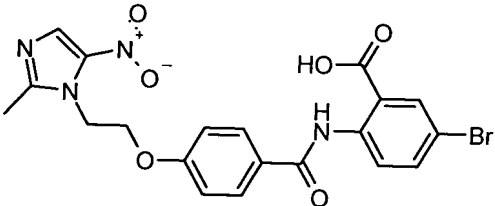
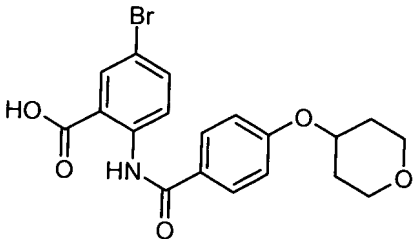
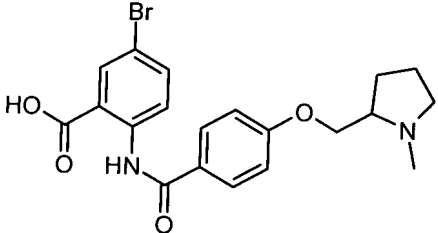
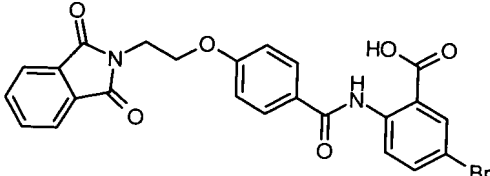
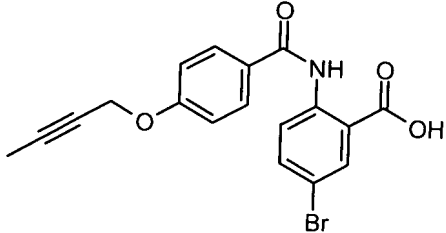
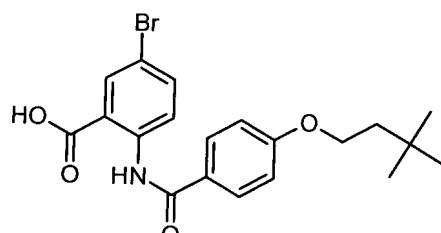
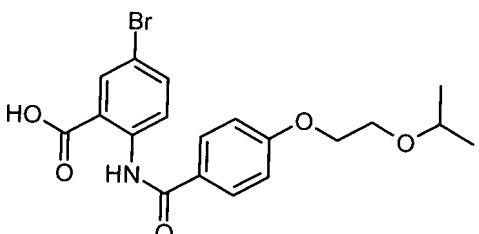
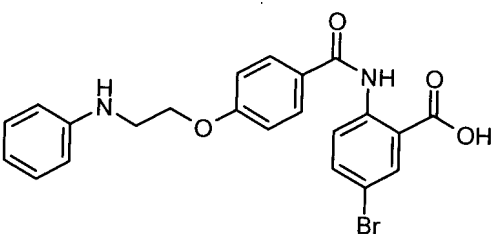
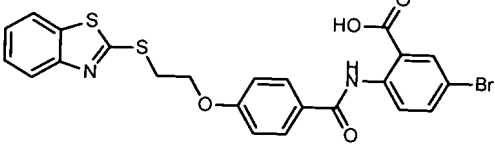
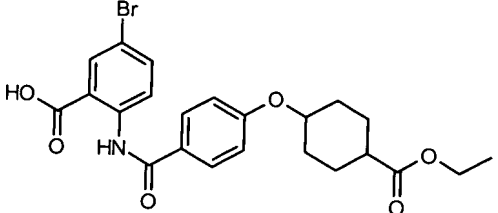
Compound No., Structure	Compound No., Structure
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PHA-563351 	PHA-563353 
PHA-563354 	PHA-563360 

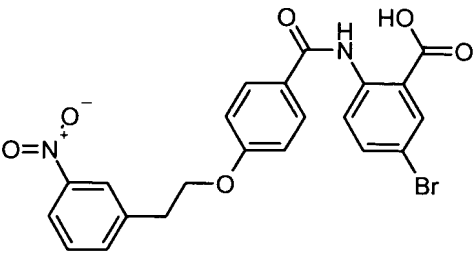
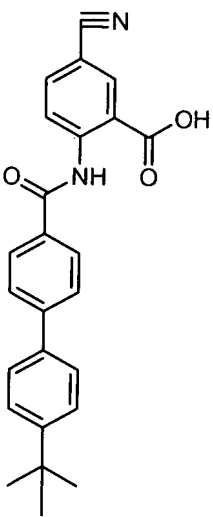
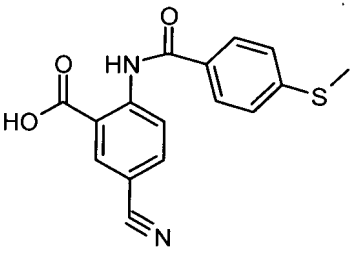
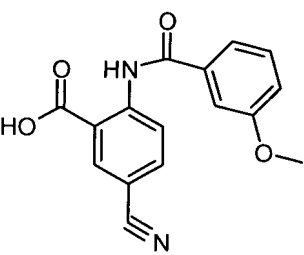
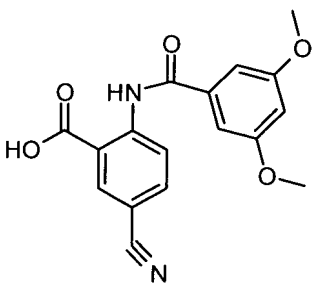
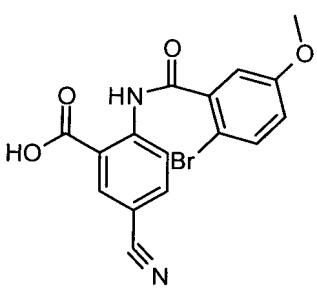
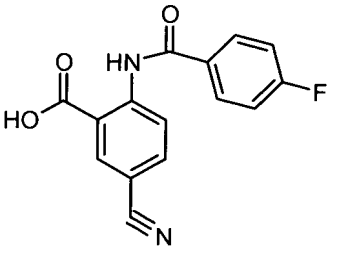
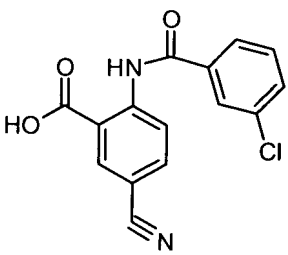
Compound No., Structure	Compound No., Structure
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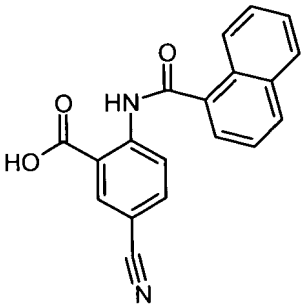
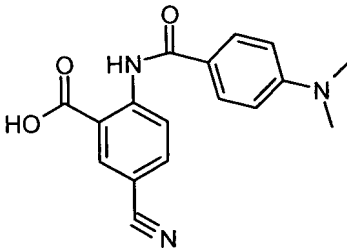
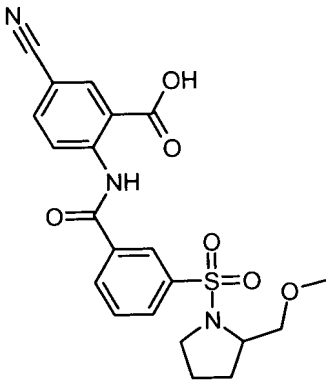
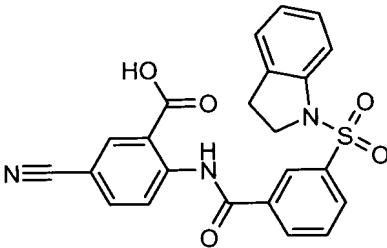
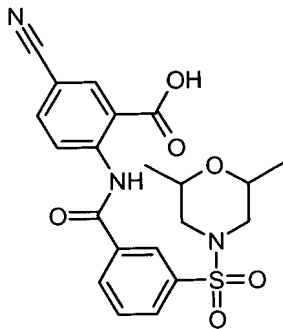
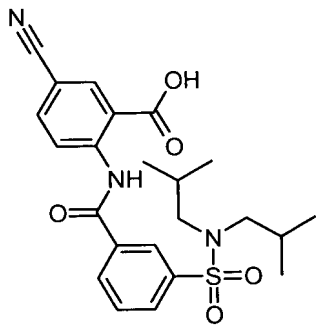
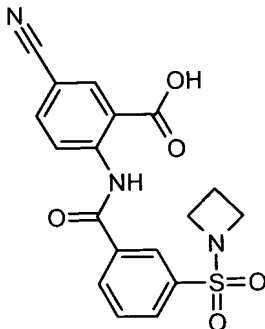
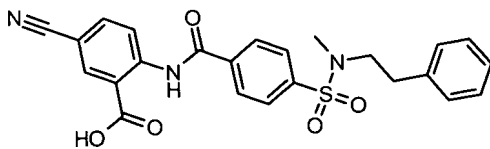
Compound No., Structure	Compound No., Structure
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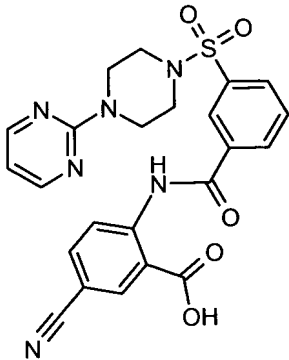
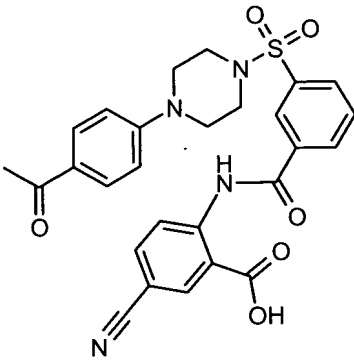
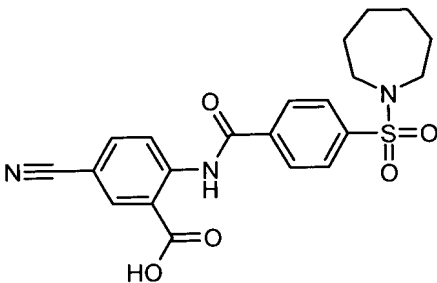
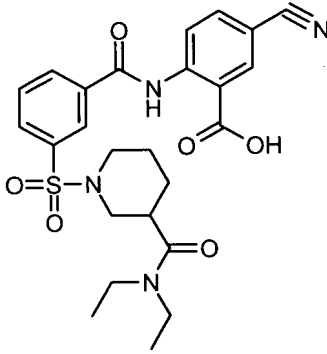
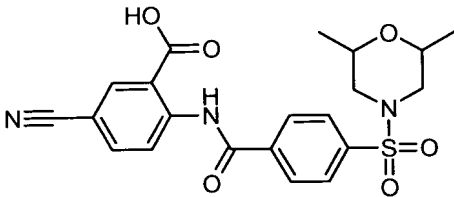
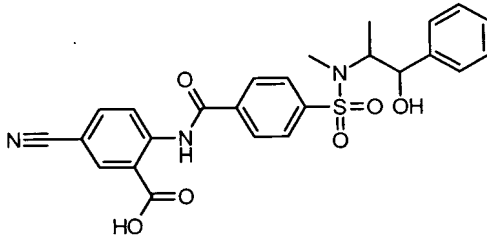
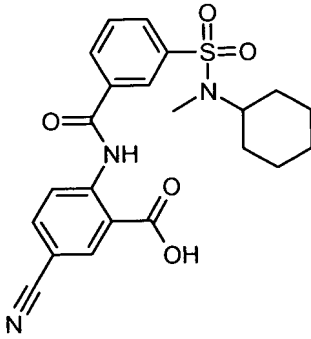
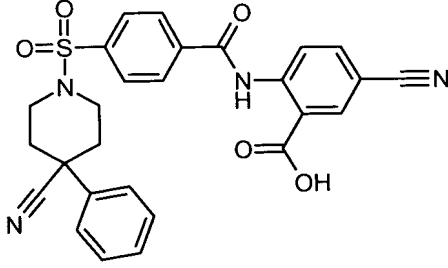
Compound No., Structure	Compound No., Structure
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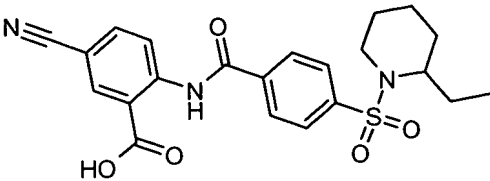
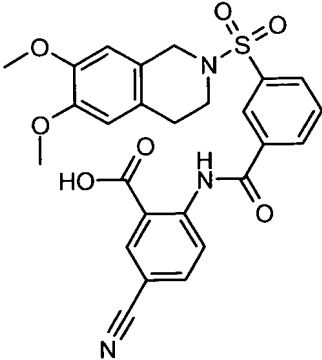
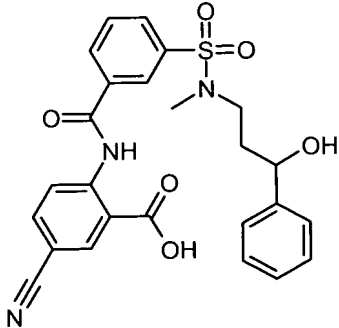
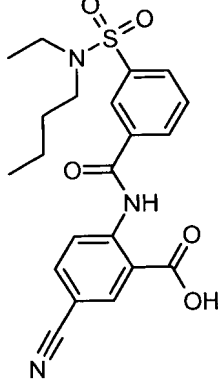
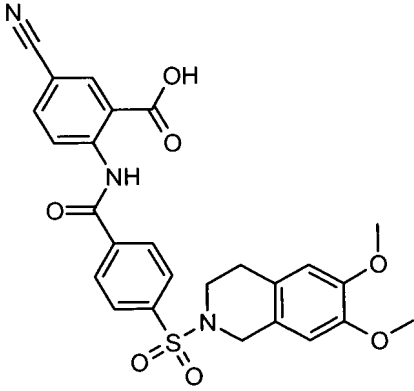
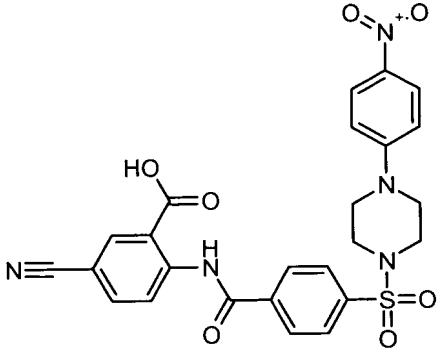


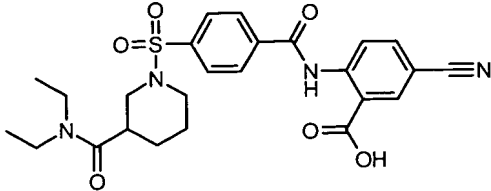
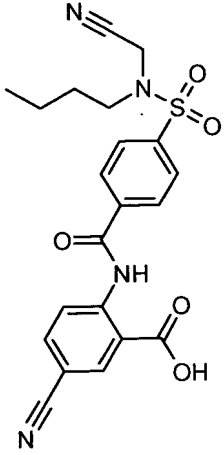
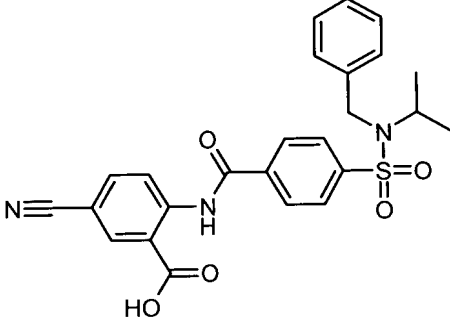
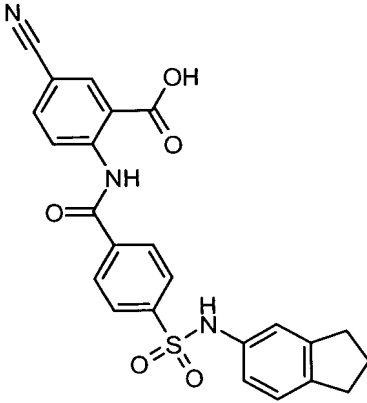
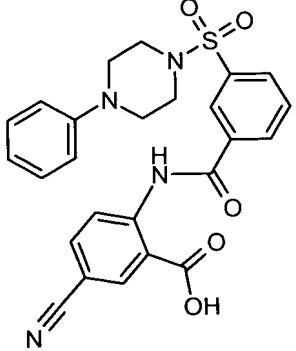
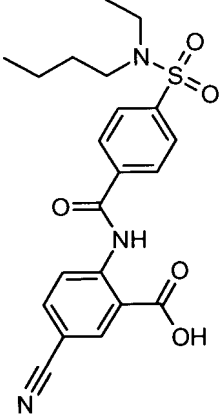
Compound No., Structure	Compound No., Structure
PHA-563413 	PHA-563415 
PHA-563417 	PHA-563419 
PHA-563420 	PHA-563426 
PHA-563427 	PHA-563440 
PHA-563441 	PHA-563442 

Compound No., Structure	Compound No., Structure
PHA-563449 	PHA-569976 
PHA-571150 	PHA-571151 
PHA-571152 	PHA-571153 
PHA-571154 	PHA-571155 

Compound No., Structure	Compound No., Structure
<p>PHA-571156</p> 	<p>PHA-571157</p> 
<p>PHA-571160</p> 	<p>PHA-571161</p> 
<p>PHA-571162</p> 	<p>PHA-571164</p> 
<p>PHA-571167</p> 	<p>PHA-571169</p> 

Compound No., Structure	Compound No., Structure
<p>PHA-571170</p> 	<p>PHA-571172</p> 
<p>PHA-571174</p> 	<p>PHA-571176</p> 
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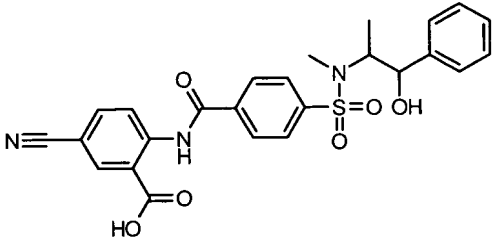
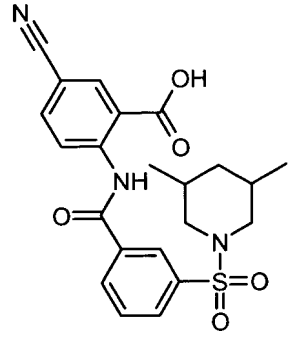
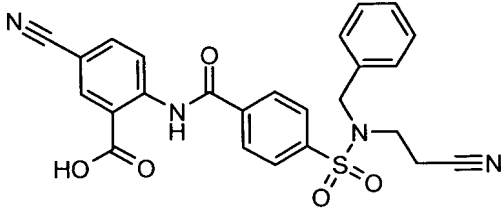
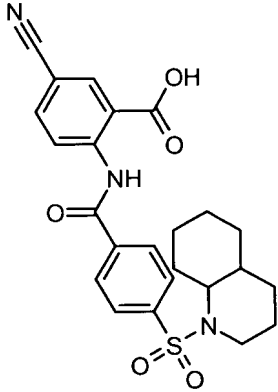
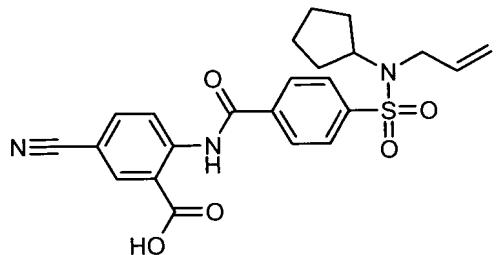
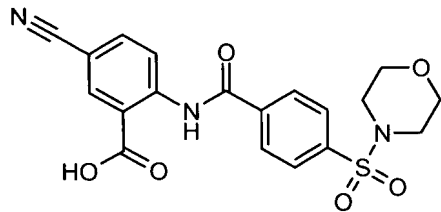
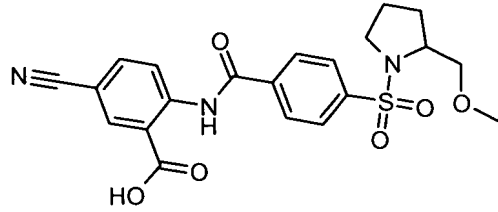
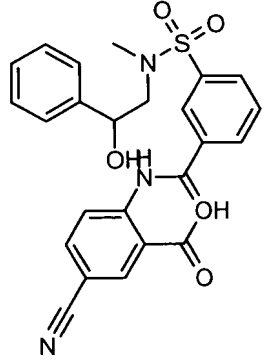
Compound No., Structure	Compound No., Structure
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<p data-bbox="266 659 440 693">PHA-571196</p> 	<p data-bbox="834 659 1008 693">PHA-571197</p> 
<p data-bbox="266 1142 440 1176">PHA-571198</p> 	<p data-bbox="834 1142 1008 1176">PHA-571199</p> 

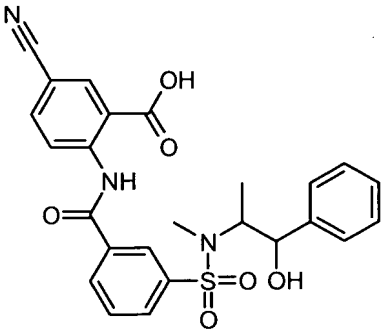
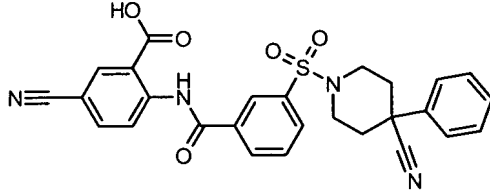
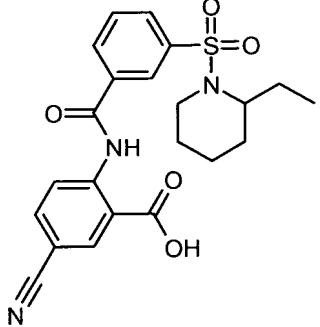
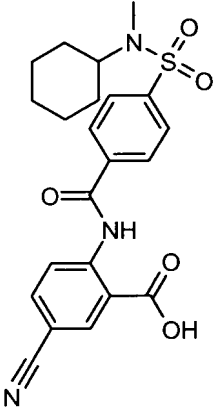
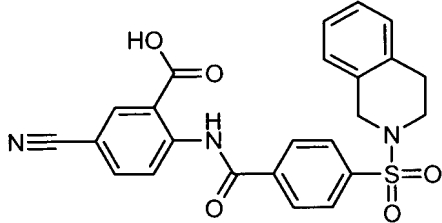
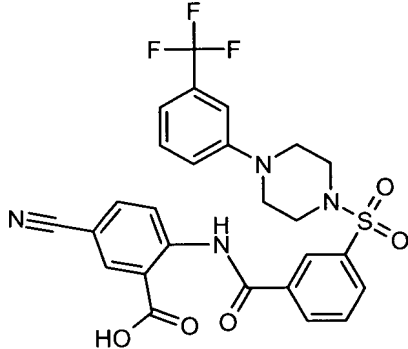
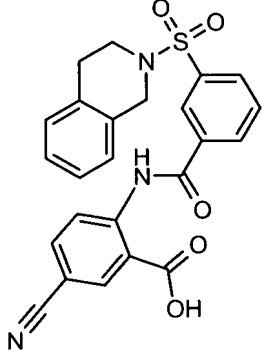
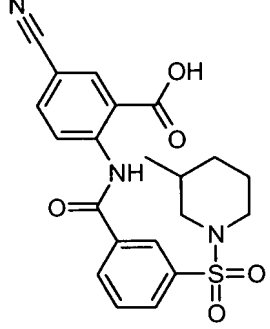
Compound No., Structure	Compound No., Structure
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<p data-bbox="264 751 446 783">PHA-571205</p> 	<p data-bbox="834 751 1016 783">PHA-571207</p> 
<p data-bbox="264 1255 451 1287">PHA-571208</p> 	<p data-bbox="834 1255 1019 1287">PHA-571214</p> 

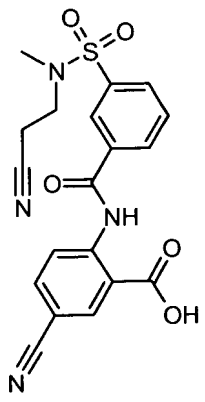
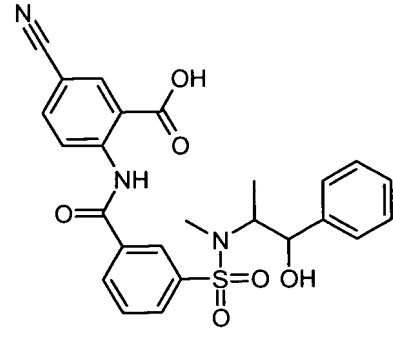
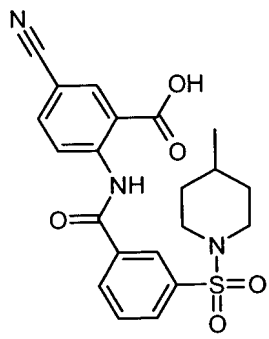
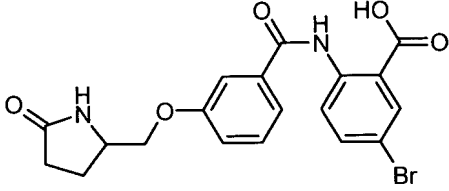
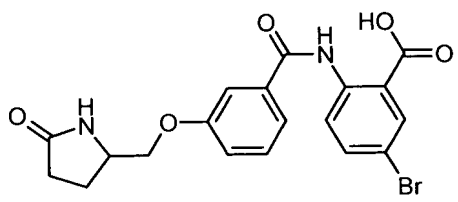
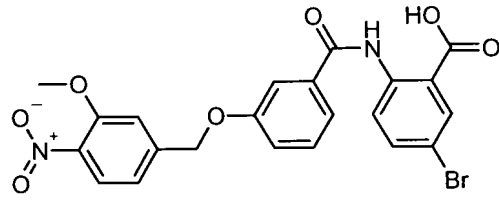
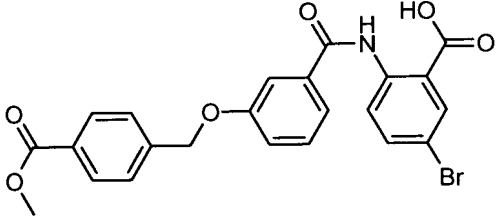
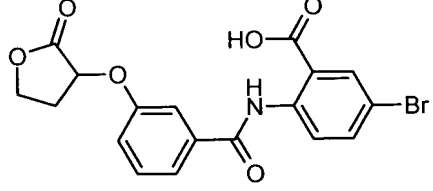
Compound No., Structure	Compound No., Structure
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PHA-571226 	PHA-571228 
PHA-571230 	PHA-571231 

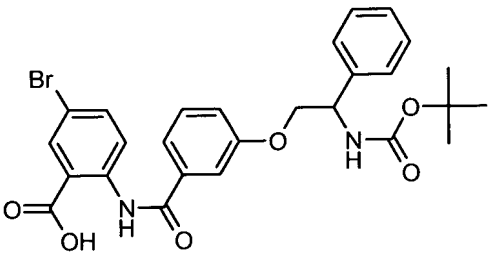
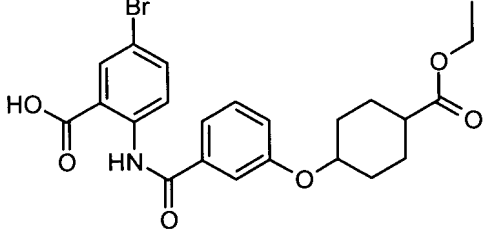
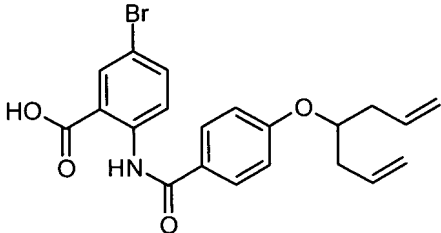
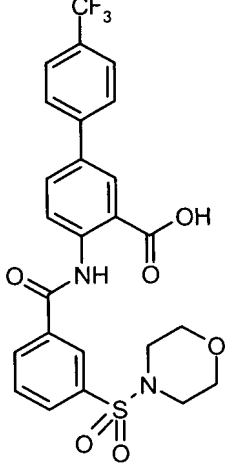
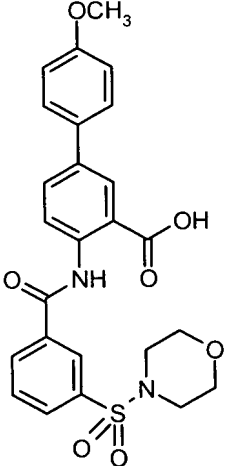
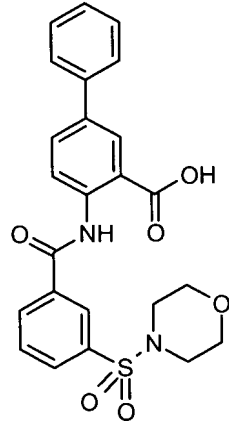
Compound No., Structure	Compound No., Structure
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PHA-571235 	PHA-571237 
PHA-571238 	PHA-571239 
PHA-571240 	PHA-571241 

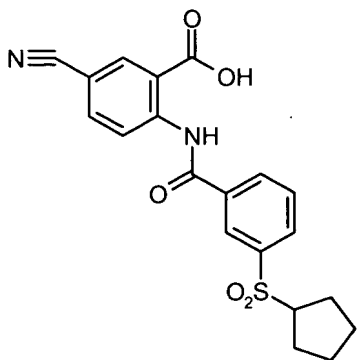
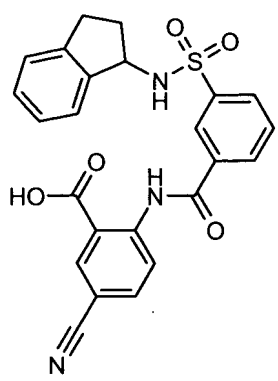
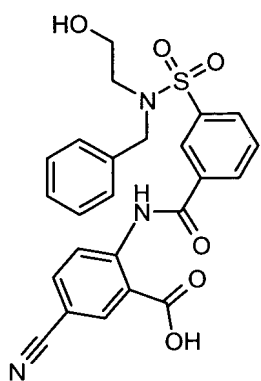
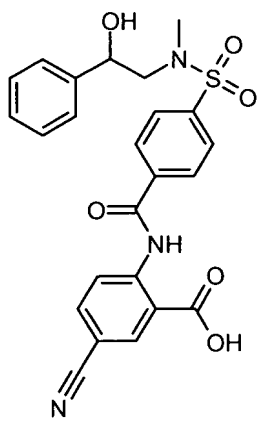
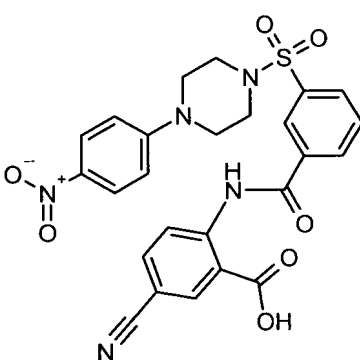
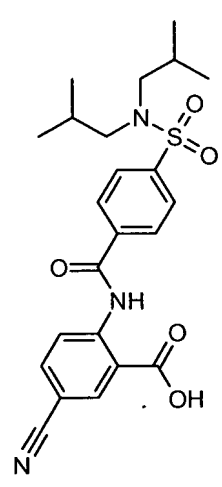


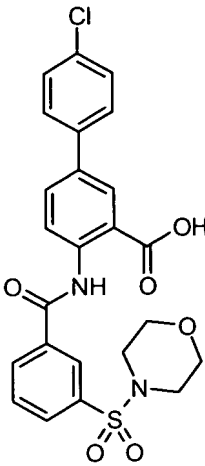
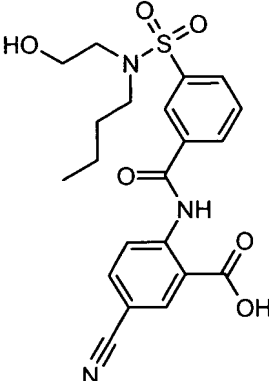
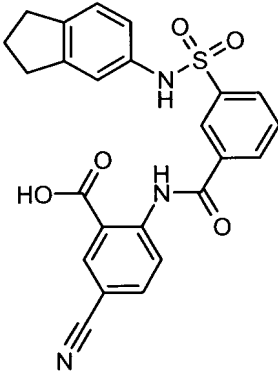
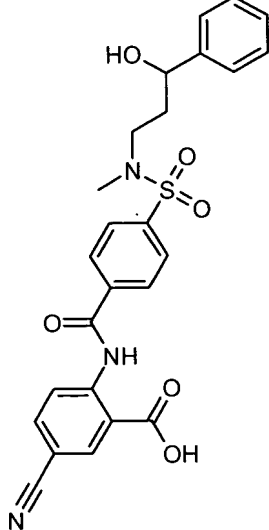
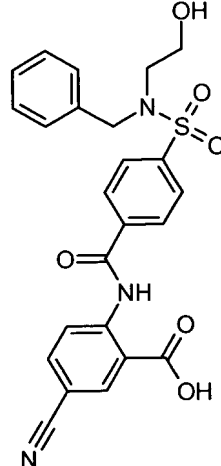
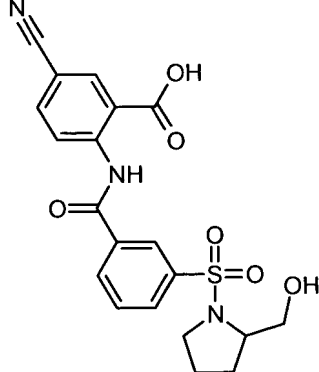
Compound No., Structure	Compound No., Structure
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<p>PHA-571246</p> 	<p>PHA-571249</p> 
<p>PHA-571253</p> 	<p>PHA-571255</p> 
<p>PHA-571257</p> 	<p>PHA-571258</p> 

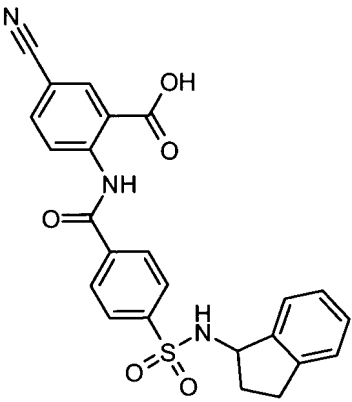
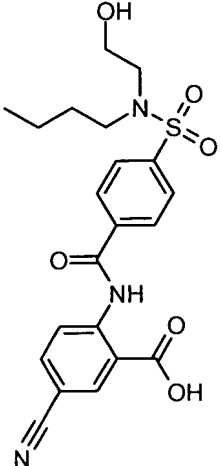
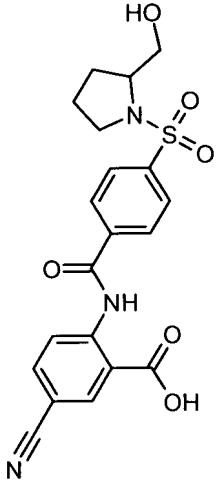
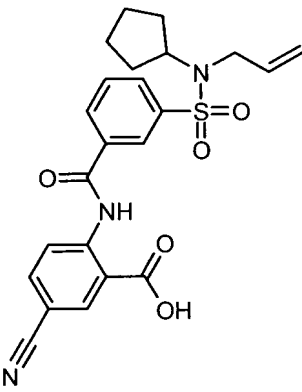
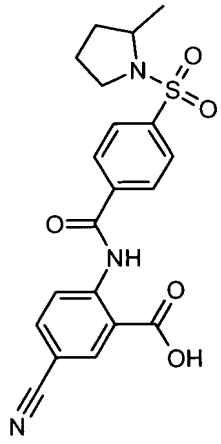
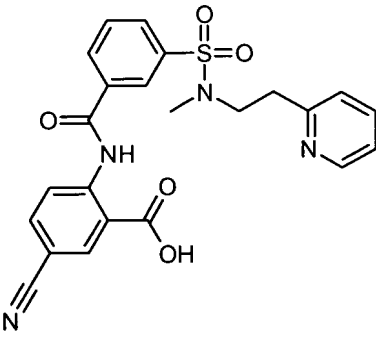
Compound No., Structure	Compound No., Structure
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<p>PHA-571263</p> 	<p>PHA-571264</p> 
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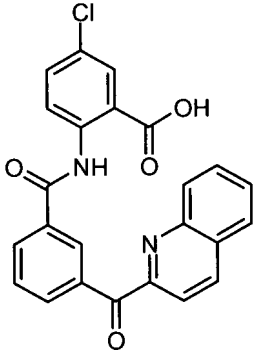
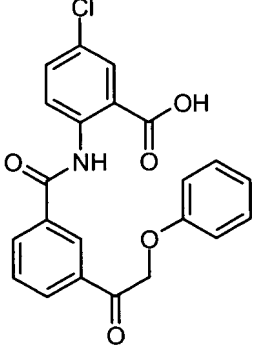
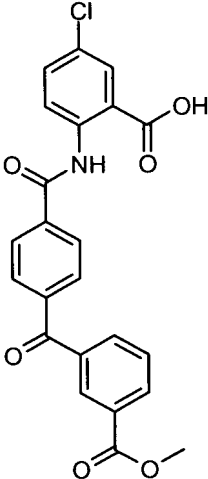
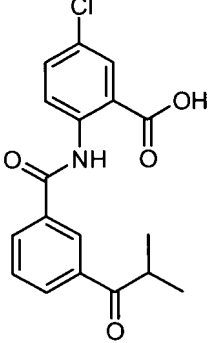
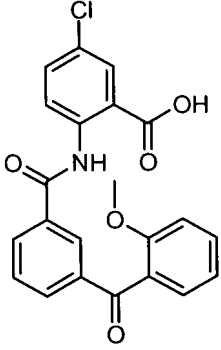
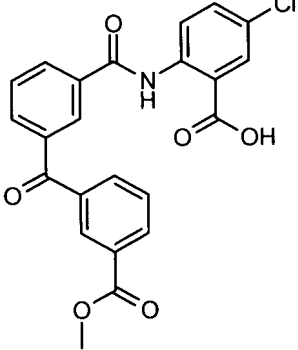
Compound No., Structure	Compound No., Structure
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<p>PHA-571273</p> 	<p>PHA-571280</p> 
<p>PHA-571281</p> 	<p>PHA-571282</p> 
<p>PHA-571283</p> 	<p>PHA-571285</p> 

Compound No., Structure	Compound No., Structure
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<p data-bbox="264 554 440 588">PHA-571292</p> 	<p data-bbox="829 554 1008 588">PHA-610940</p> 
<p data-bbox="264 1121 440 1155">PHA-610941</p> 	<p data-bbox="829 1121 1008 1155">PHA-610942</p> 

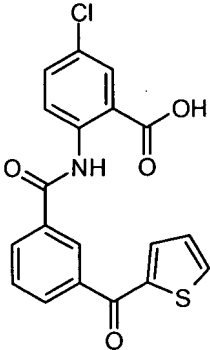
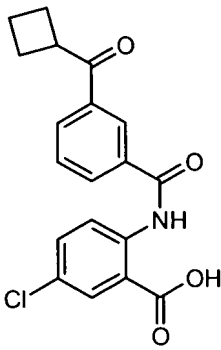
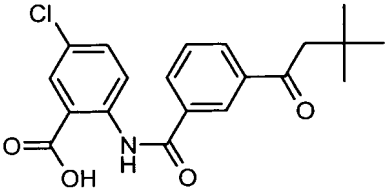
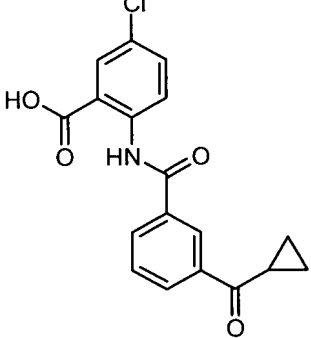
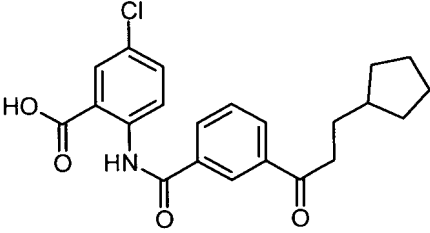
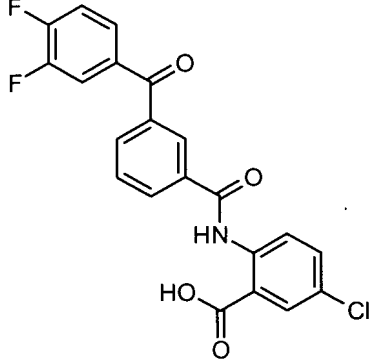
Compound No., Structure	Compound No., Structure
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PHA-656808 	PHA-656809 
PHA-656810 	PHA-656811 

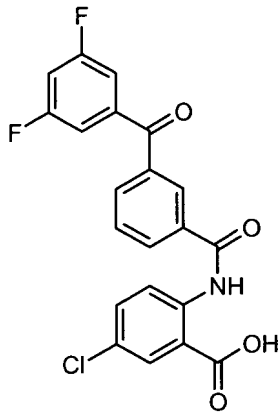
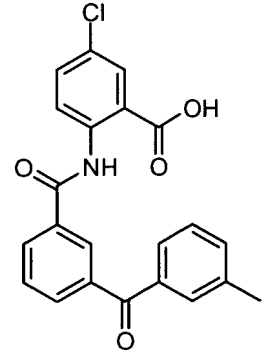
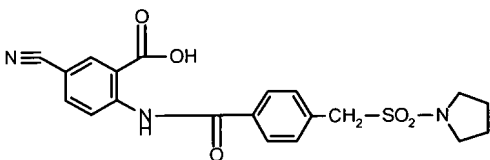
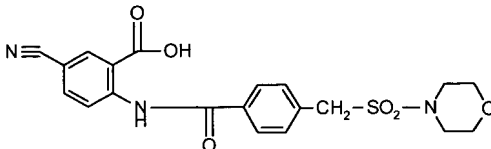
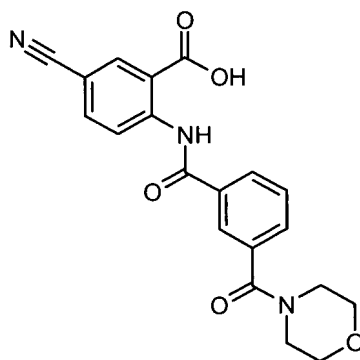
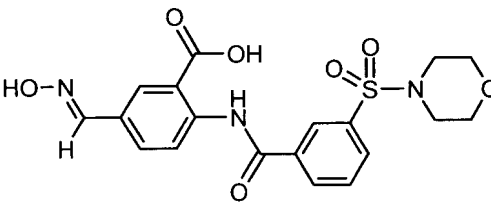
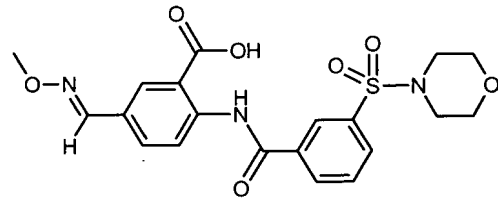
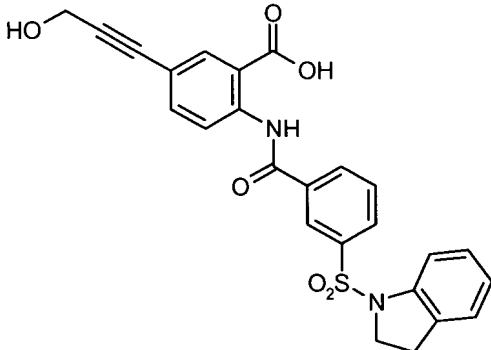
Compound No., Structure	Compound No., Structure
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<p data-bbox="269 758 448 789">PHA-656860</p> 	<p data-bbox="837 758 1016 789">PHA-656861</p> 
<p data-bbox="269 1398 448 1430">PHA-656862</p> 	<p data-bbox="837 1398 1016 1430">PHA-656863</p> 

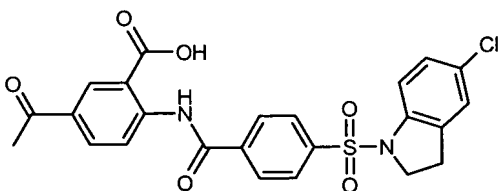
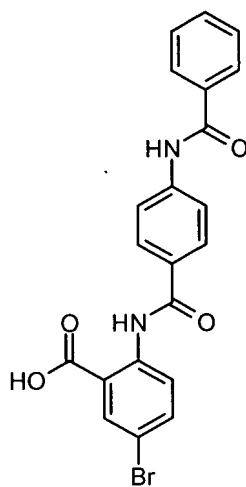
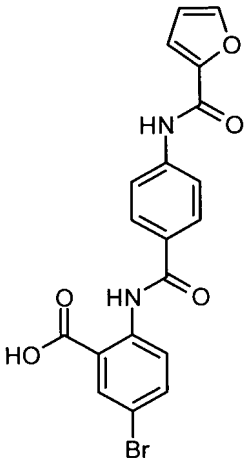
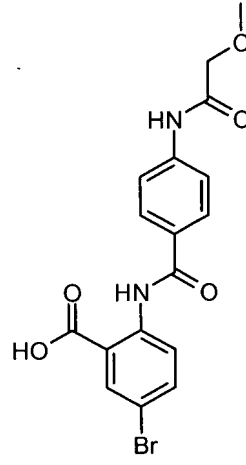
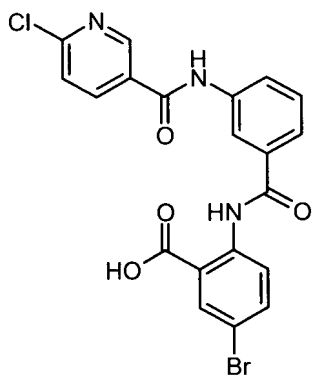
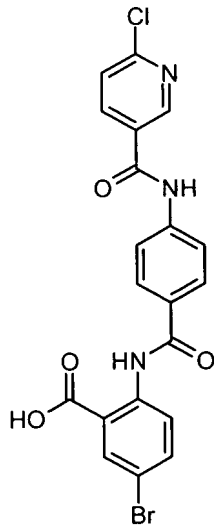
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<p data-bbox="272 772 446 804">PHA-656868</p> 	<p data-bbox="841 772 1015 804">PHA-656870</p> 
<p data-bbox="272 1350 446 1381">PHA-656871</p> 	<p data-bbox="841 1350 1015 1381">PHA-656872</p> 

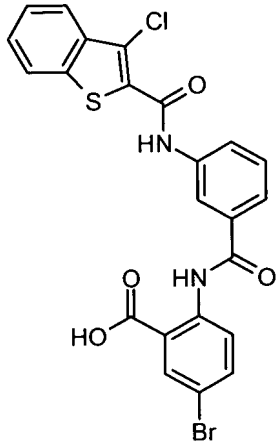
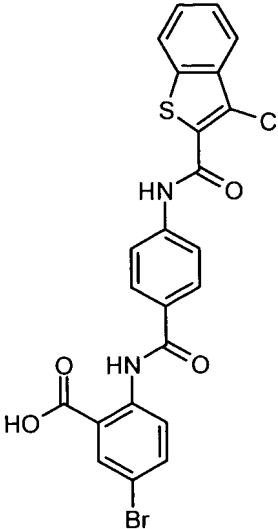
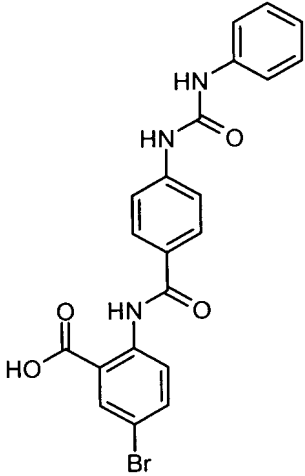
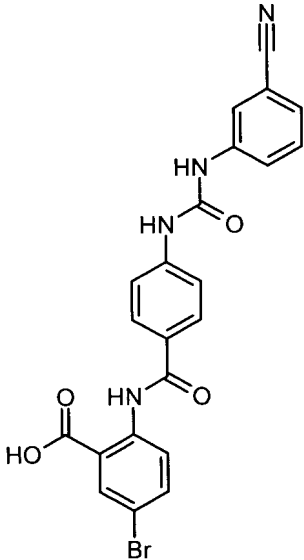
Compound No., Structure	Compound No., Structure
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<p data-bbox="267 657 446 688">PHA-656883</p> 	<p data-bbox="836 657 1015 688">PHA-656884</p> 
<p data-bbox="267 1234 446 1266">PHA-656885</p> 	<p data-bbox="836 1234 1015 1266">PHA-656886</p> 

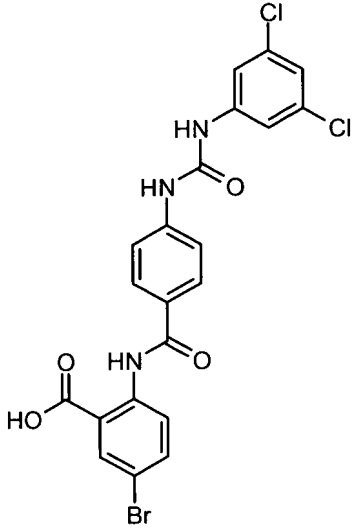
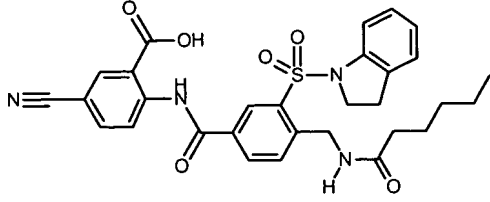
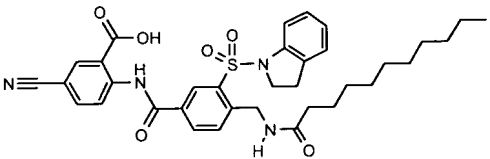
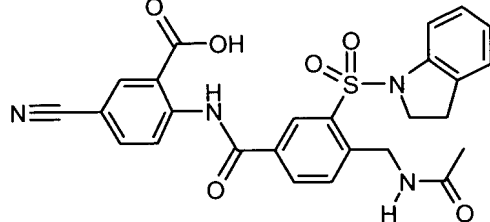
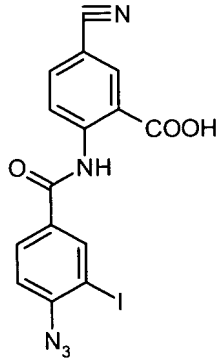
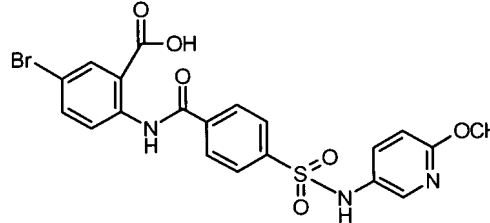


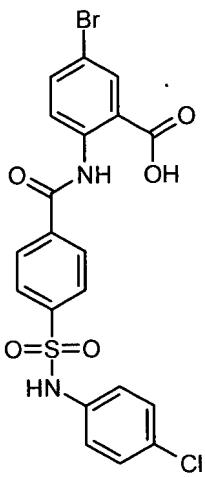
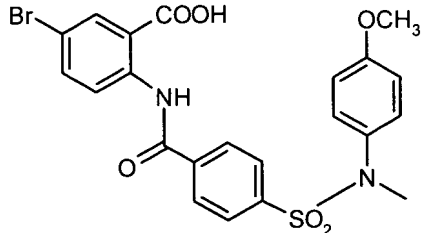
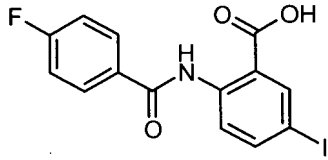
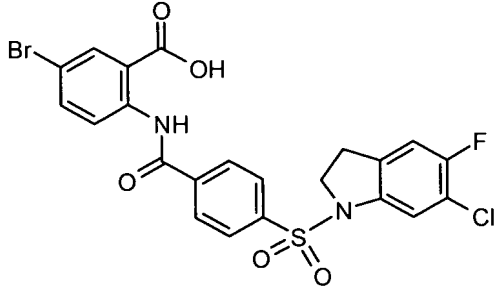
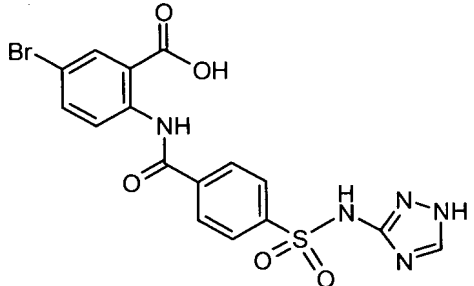
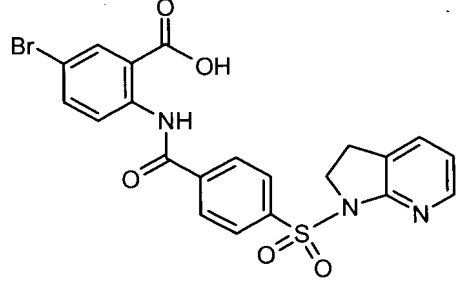
Compound No., Structure	Compound No., Structure
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<p data-bbox="269 657 448 688">PHA-656889</p>  <chem data-bbox="350 720 732 909">Clc1ccc(cc1C(=O)O)NC(=O)c2ccccc2C(=O)CC(C)(C)C</chem>	<p data-bbox="837 657 1016 688">PHA-656890</p>  <chem data-bbox="954 720 1263 1056">Clc1ccc(cc1C(=O)O)NC(=O)c2ccccc2C(=O)C1CC1</chem>
<p data-bbox="269 1098 448 1129">PHA-656891</p>  <chem data-bbox="329 1161 756 1392">Clc1ccc(cc1C(=O)O)NC(=O)c2ccccc2C(=O)CC1CCCC1</chem>	<p data-bbox="837 1098 1016 1129">PHA-656892</p>  <chem data-bbox="930 1161 1300 1518">Fc1cc(F)ccc1C(=O)c2ccc(cc2NC(=O)c3cc(Cl)ccc3C(=O)O)c4ccccc4</chem>

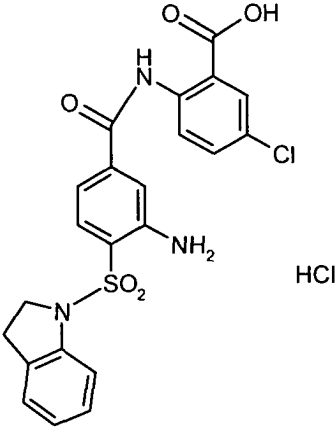
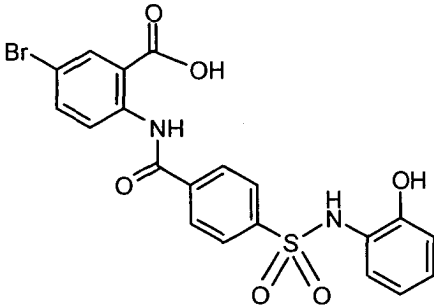
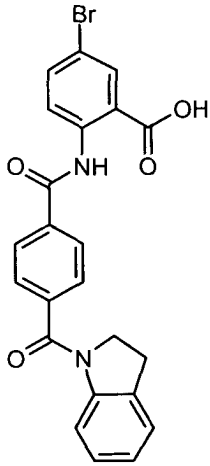
Compound No., Structure	Compound No., Structure
<p>PHA-656893</p> 	<p>PHA-656894</p> 
<p>PHA-662253</p> 	<p>PHA-662254</p> 
<p>PHA-662412</p> 	<p>PHA-679756</p> 
<p>PHA-679759</p> 	<p>PHA-687570</p> 

Compound No., Structure	Compound No., Structure
<p data-bbox="267 205 446 237">PHA-708922</p> 	<p data-bbox="836 205 1015 237">PHA-708977</p> 
<p data-bbox="267 789 446 821">PHA-708979</p> 	<p data-bbox="836 789 1015 821">PHA-708987</p> 
<p data-bbox="267 1350 446 1381">PHA-713389</p> 	<p data-bbox="836 1350 1015 1381">PHA-713390</p> 

Compound No., Structure	Compound No., Structure
<p data-bbox="264 207 440 241">PHA-713391</p>  <p>The structure of PHA-713391 consists of a 5-chloro-1H-benzothiophene-2-carboxamide group linked via its amide nitrogen to a para-phenylene ring. This para-phenylene ring is further linked via its other amide nitrogen to a meta-phenylene ring, which also bears a carboxylic acid group and a bromine atom at the 4-position.</p>	<p data-bbox="834 207 1010 241">PHA-713392</p>  <p>The structure of PHA-713392 is similar to PHA-713391, but the meta-phenylene ring at the bottom is replaced by a para-phenylene ring. The rest of the molecule, including the 5-chloro-1H-benzothiophene-2-carboxamide group and the intermediate para-phenylene ring, remains the same.</p>
<p data-bbox="264 842 446 875">PHA-713393</p>  <p>The structure of PHA-713393 features a 4-phenylbenzamide group linked via its amide nitrogen to a para-phenylene ring. This para-phenylene ring is further linked via its other amide nitrogen to a meta-phenylene ring, which also bears a carboxylic acid group and a bromine atom at the 4-position.</p>	<p data-bbox="834 842 1016 875">PHA-713395</p>  <p>The structure of PHA-713395 is similar to PHA-713393, but the 4-phenylbenzamide group is replaced by a 4-cyanophenylbenzamide group, where the phenyl ring at the top has a cyano group at the para position.</p>

Compound No., Structure	Compound No., Structure
<p data-bbox="277 212 448 239">PHA-713397</p> 	<p data-bbox="839 212 1010 239">PHA-738531</p> 
<p data-bbox="277 829 448 856">PHA-738532</p> 	<p data-bbox="839 829 1010 856">PHA-740499</p> 
<p data-bbox="277 1157 448 1184">PHA-748361</p> 	<p data-bbox="839 1157 1010 1184">PNU-276556</p> 

Compound No., Structure	Compound No., Structure
<p>PNU-276672</p> 	<p>PNU-276873</p> 
<p>PNU-281164</p> 	<p>PNU-282858</p> 
<p>PNU-282859</p> 	<p>PNU-282860</p> 

Compound No., Structure	Compound No., Structure
<p>PNU-290881A</p> 	<p>PNU-291997</p> 
<p>PNU-292577</p> 	

### Example 11: ACTIVITY DATA

#### MIC Test Method

The *in vitro* MICs of test compounds were determined by a standard agar dilution method. A stock drug solution of each analog was prepared in the preferred solvent, usually DMSO:H<sub>2</sub>O (1:3). Serial 2-fold dilutions of each sample are made using 1.0 ml aliquots of sterile distilled water. To each 1.0 ml aliquot of drug was added 9 ml of molten Mueller Hinton agar medium. The drug-supplemented agar was mixed, poured into 15 x 100 mm petri dishes, and allowed to solidify and dry prior to inoculation.

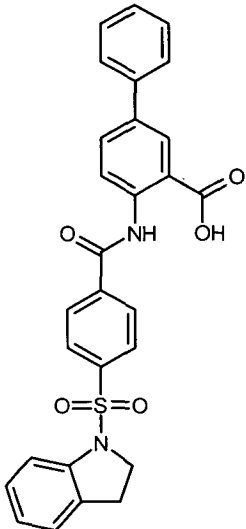
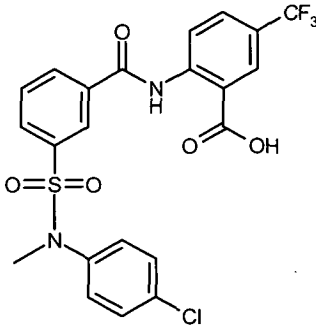
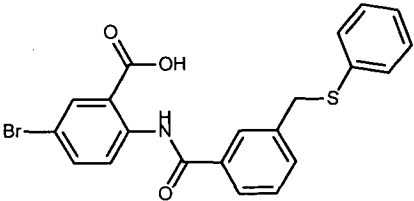
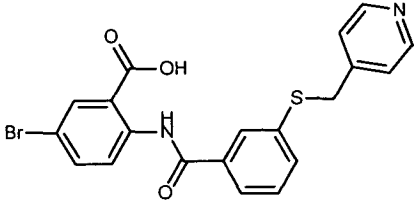
Vials of each of the test organisms are maintained frozen in the vapor phase of a liquid nitrogen freezer. Test cultures are grown overnight at 35°C on the medium

appropriate for the organism. Colonies are harvested with a sterile swab, and cell suspensions are prepared in Trypticase Soy broth (TSB) to equal the turbidity of a 0.5 McFarland standard. A 1:20 dilution of each suspension was made in TSB. The plates containing the drug supplemented agar are inoculated with a 0.001 ml drop of the cell suspension using a Steers replicator, yielding approximately  $10^4$  to  $10^5$  cells per spot. The plates are incubated overnight at 35°C.

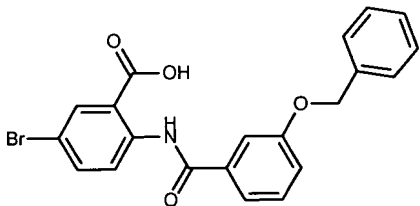
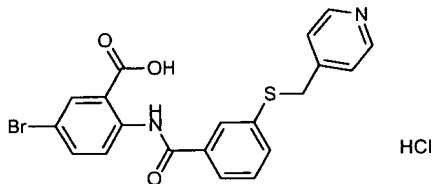
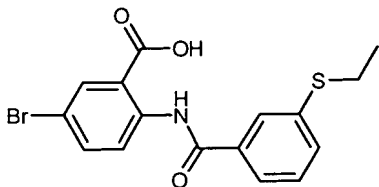
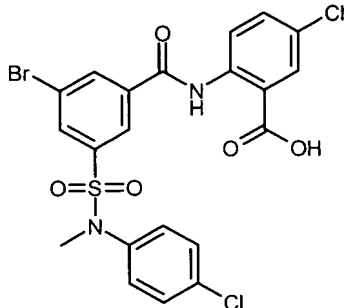
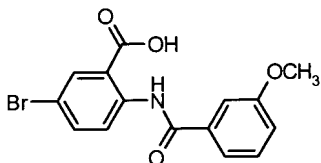
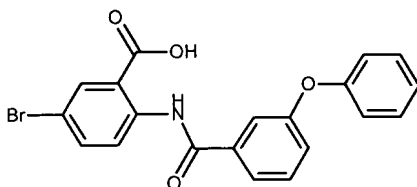
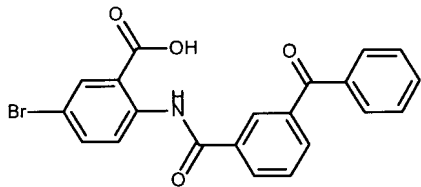
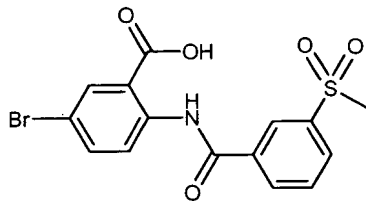
Following incubation the Minimum Inhibitory Concentration (MIC  $\mu\text{g/ml}$ ), the lowest concentration of drug that inhibits visible growth of the organism, was read and recorded. The data is shown in Tables I and II.

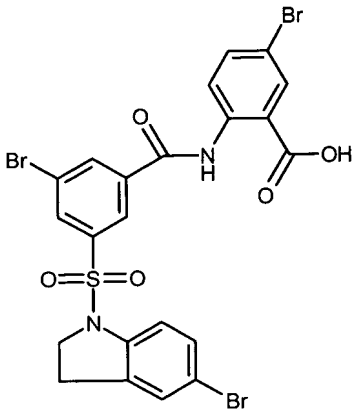
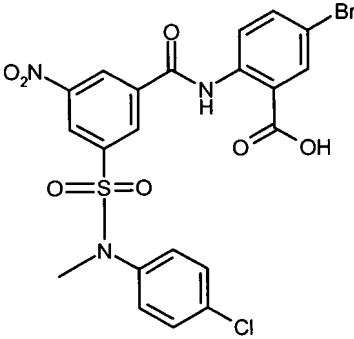
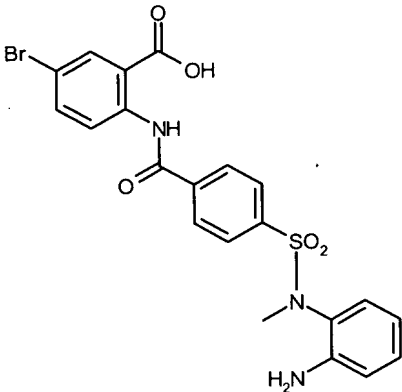
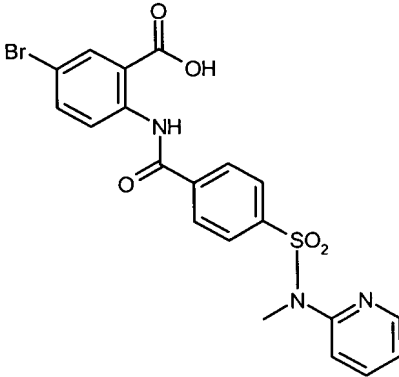
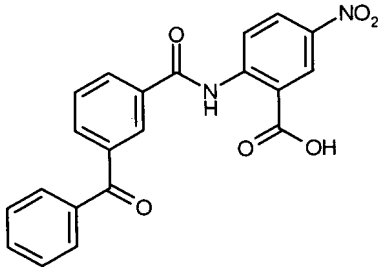
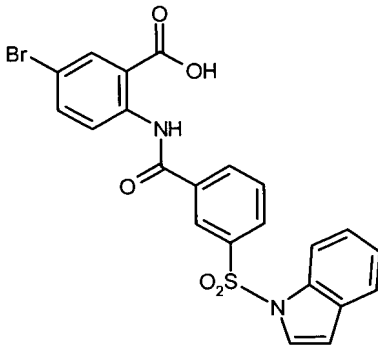
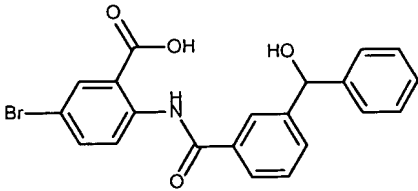
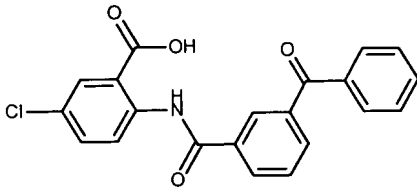
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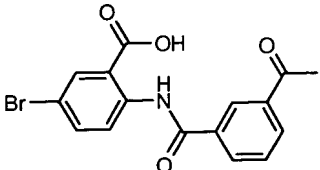
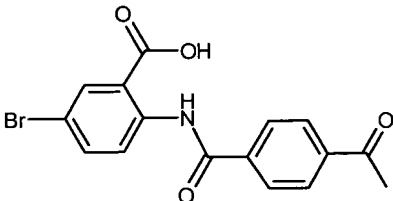
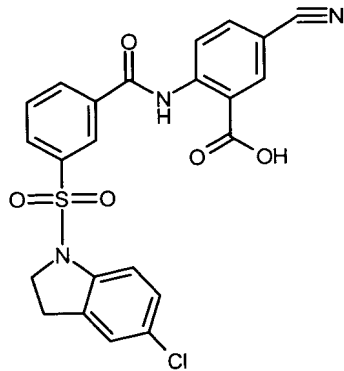
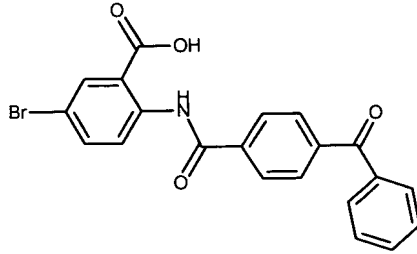
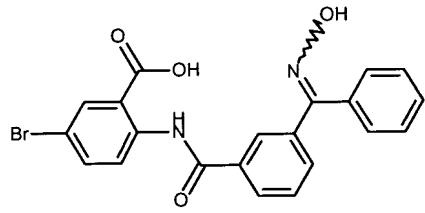
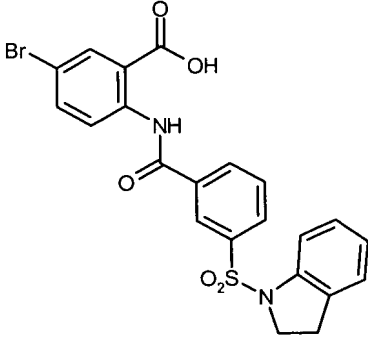
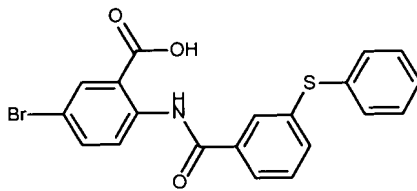
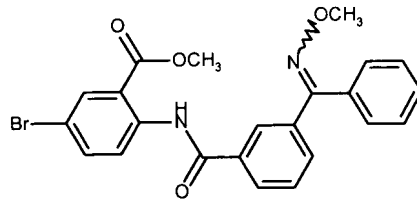
**Table 1: Activity Data**

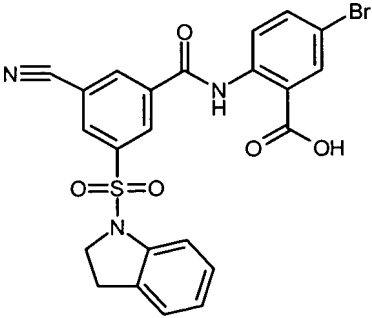
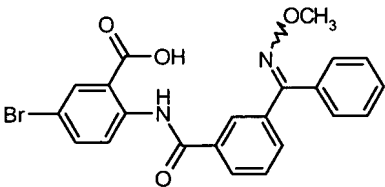
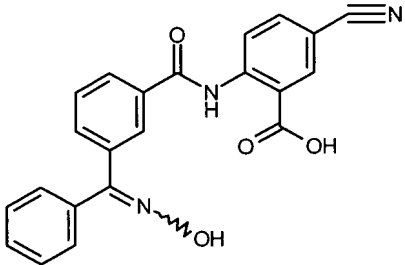
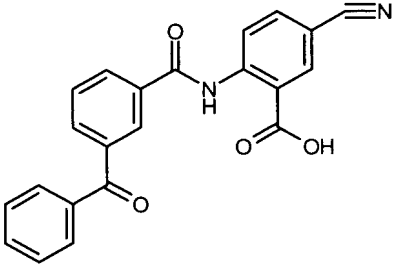
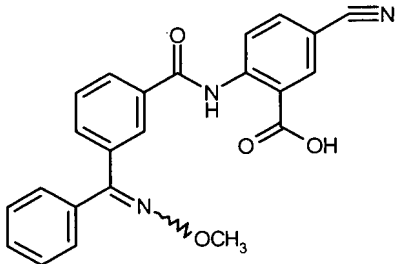
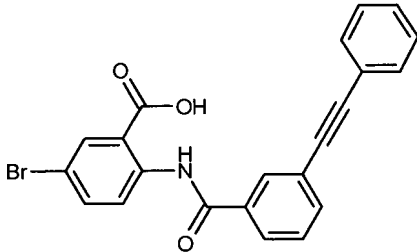
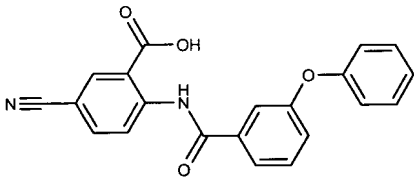
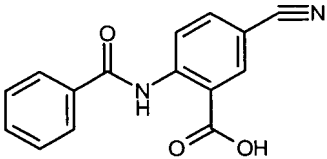
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
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PHA-501684 	1	PHA-502339 	2

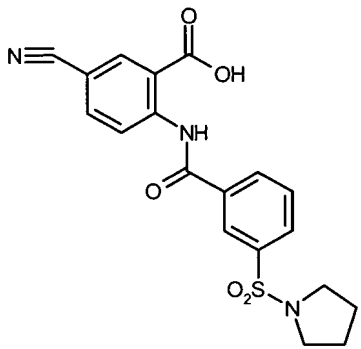
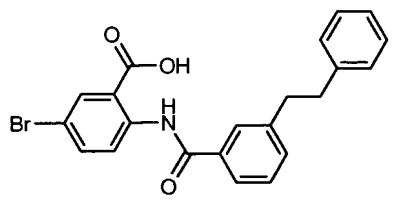
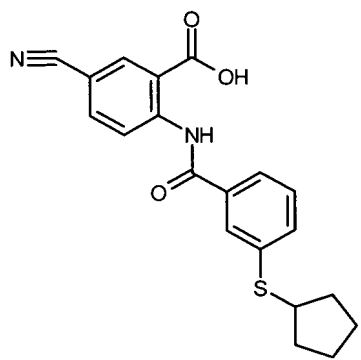
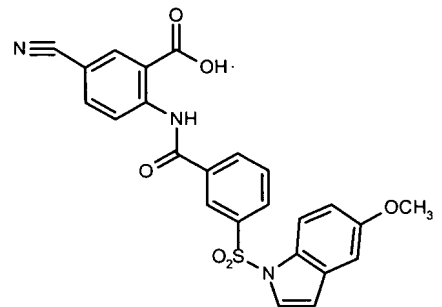
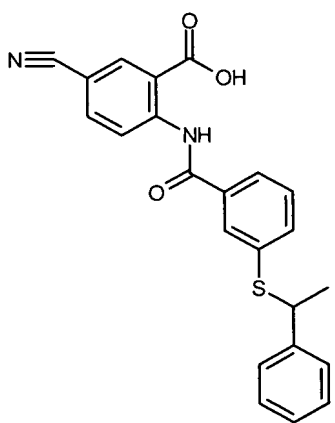
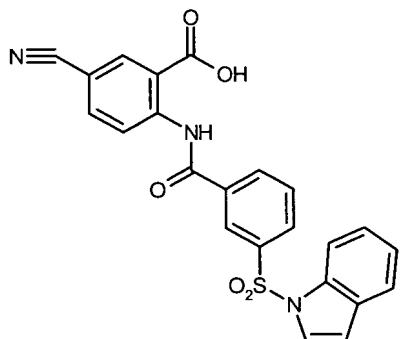


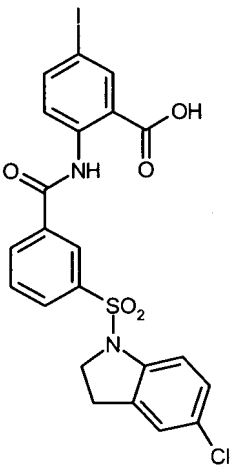
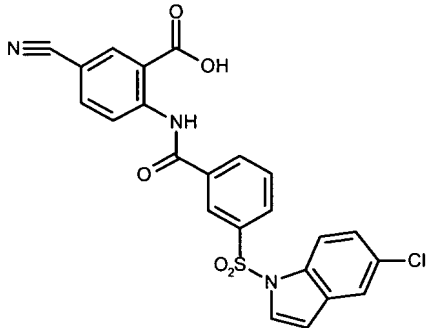
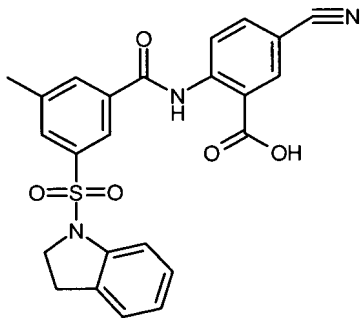
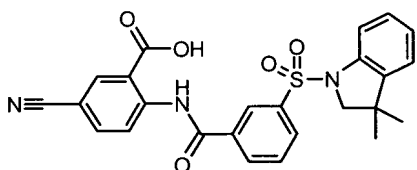
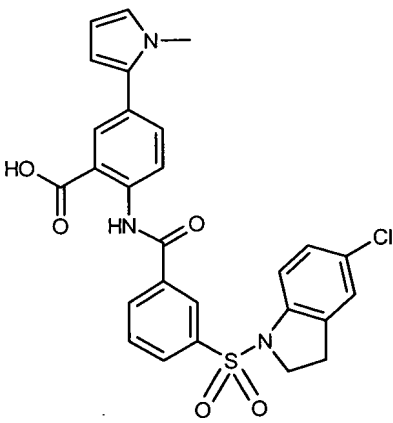
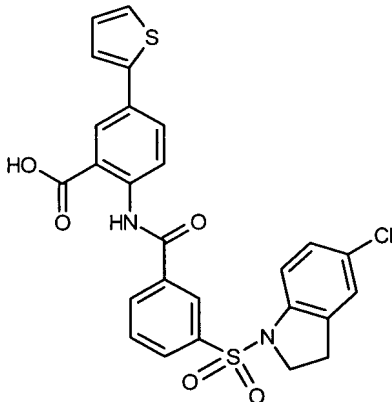
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-501685 	2	PHA-502339A 	8
PHA-501748 	2	PHA-509059 	0.5
PHA-504639 	4	PHA-513535 	2
PHA-515448 	2	PHA-513541 	64

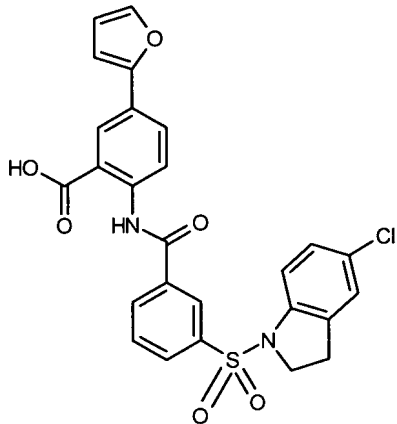
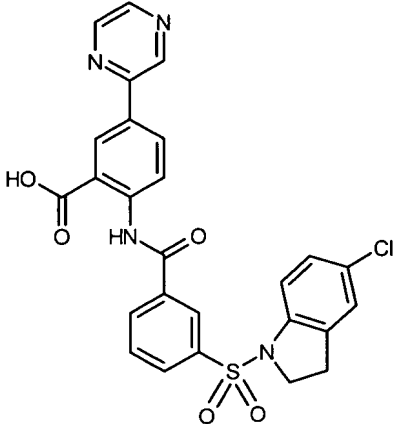
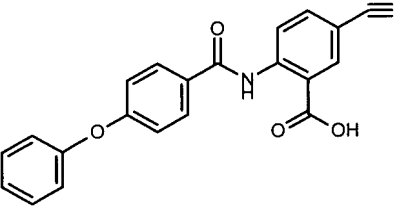
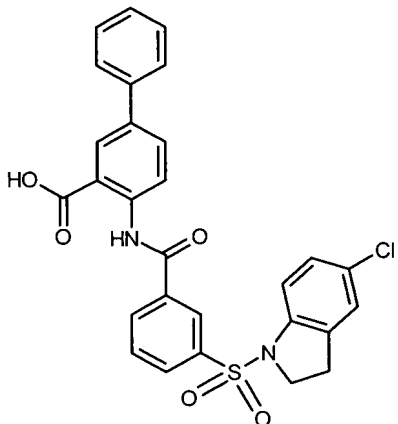
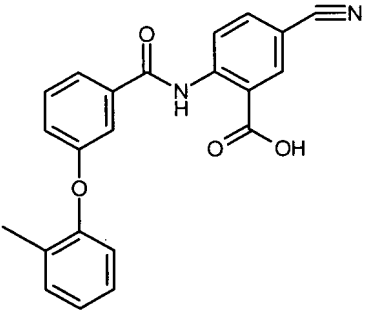
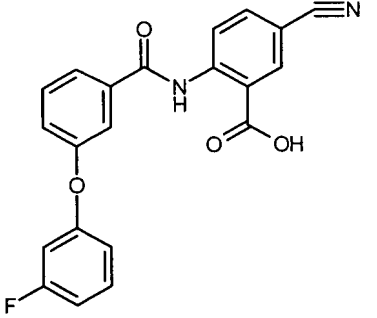
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-515585 	1	PHA-515583 	8
PHA-516113 	2	PHA-516112 	8
PHA-519402 	0.5	PHA-516116 	0.5
PHA-521534 	1	PHA-518226 	2

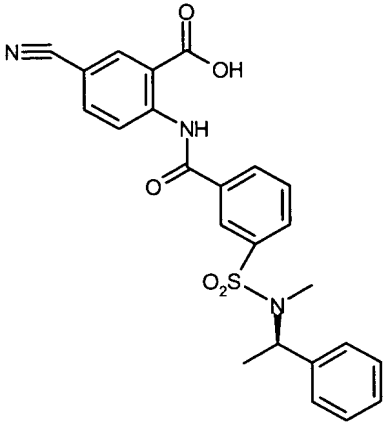
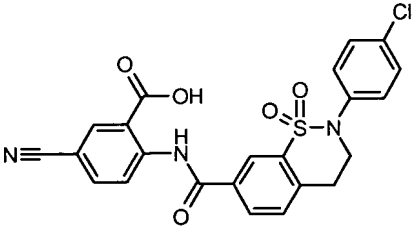
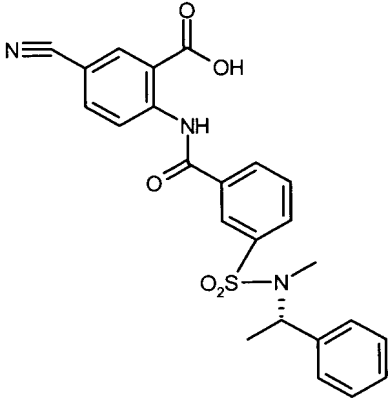
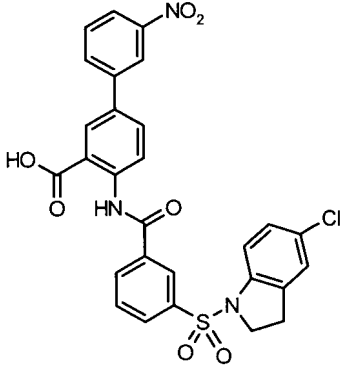
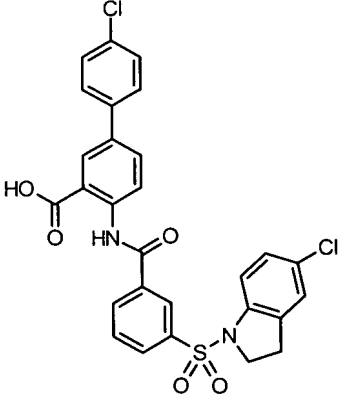
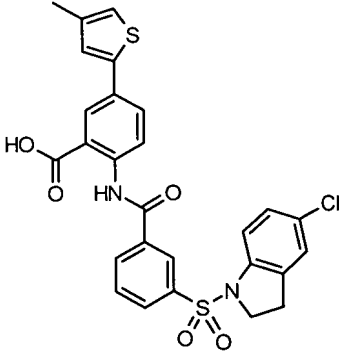
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-522145 	32	PHA-520446 	16
PHA-524523 	0.125	PHA-520447 	1
PHA-524545 	0.25	PHA-520938 	1
PHA-526580 	1	PHA-521535 	>128

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-530687 	8	PHA-522146 	0.5
PHA-535548 	0.25	PHA-524524 	1
PHA-535549 	0.25	PHA-526578 	2
PHA-535553 	1	PHA-530685 	32

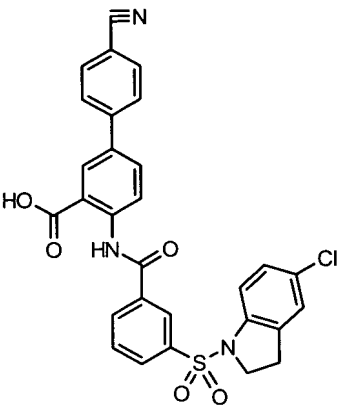
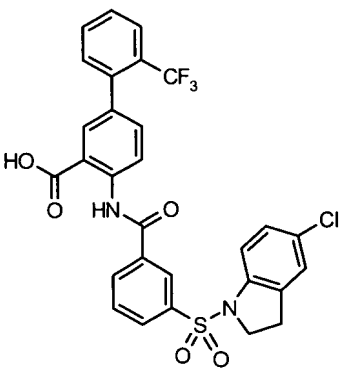
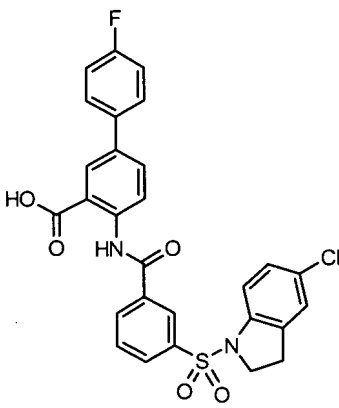
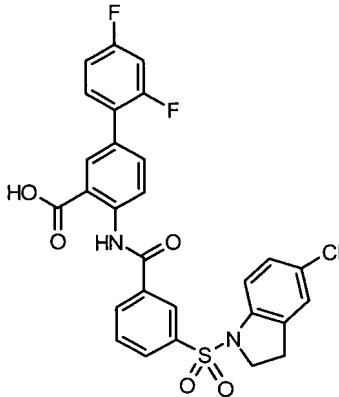
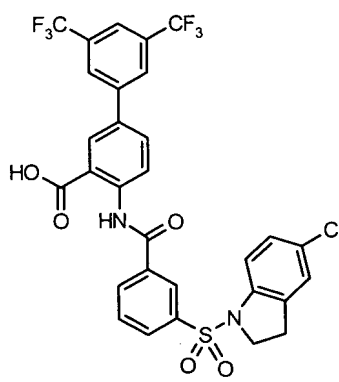
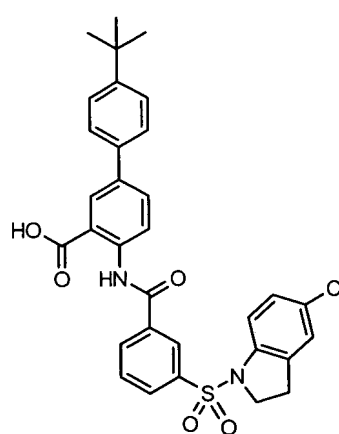
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-543140 	1	PHA-530989 	4
PHA-546926 	0.5	PHA-543139 	0.125
PHA-547267 	0.125	PHA-543141 	0.125

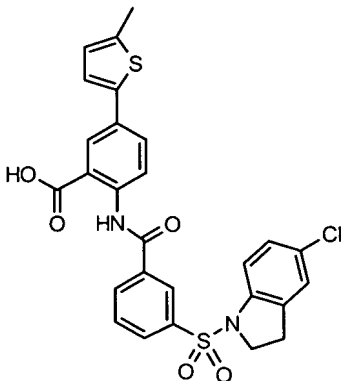
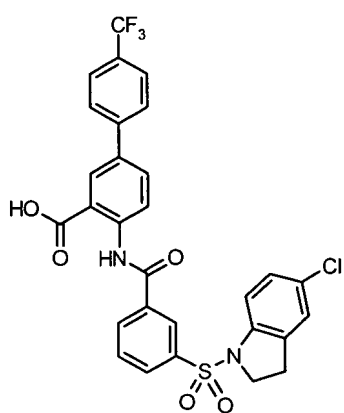
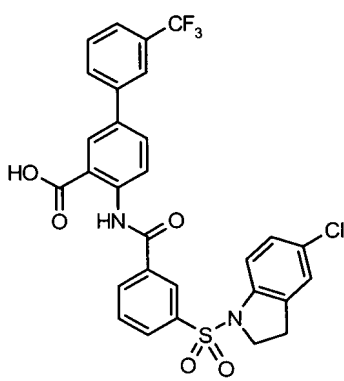
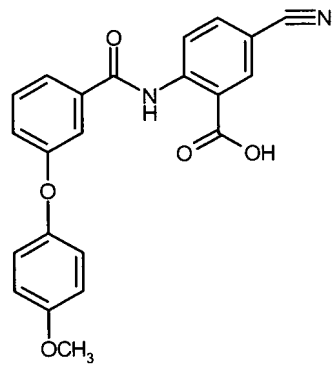
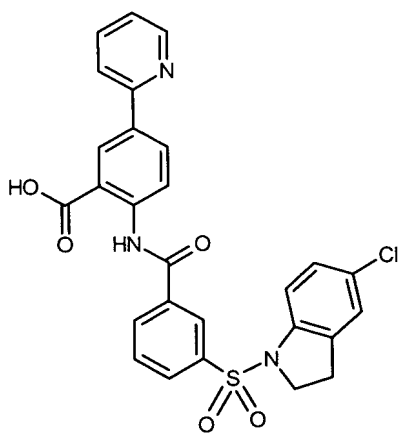
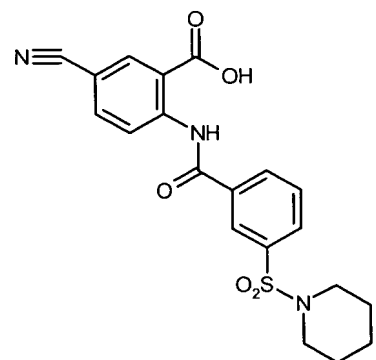
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-552831 	1	PHA-543681 	0.125
PHA-556214 	1	PHA-555027 	1
PHA-556658 	8	PHA-556657 	2

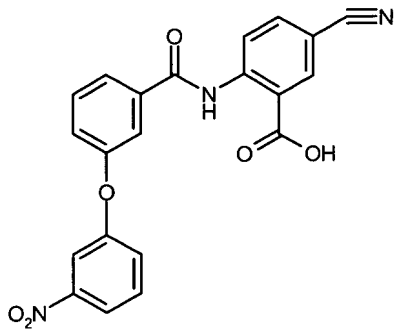
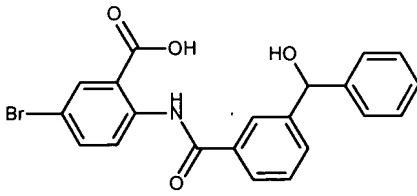
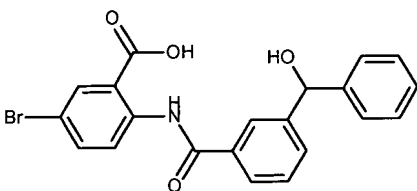
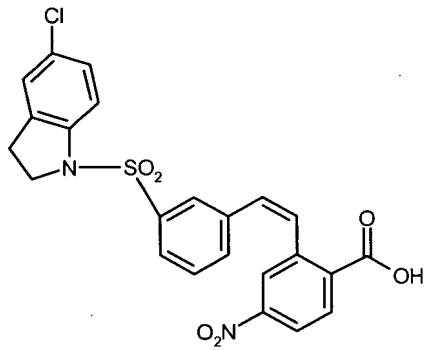
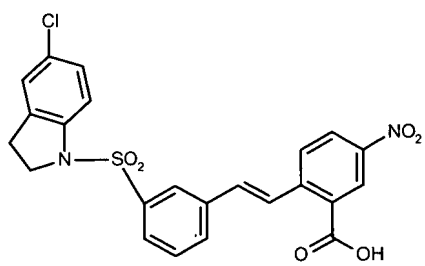
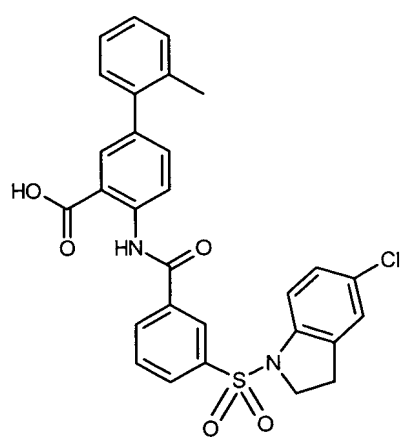
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-556663 	8	PHA-556661 	8
PHA-561055 	1	PHA-557035 	4
PHA-562733 	0.25	PHA-562731 	1

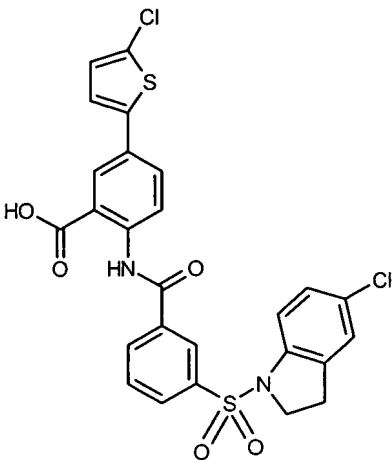
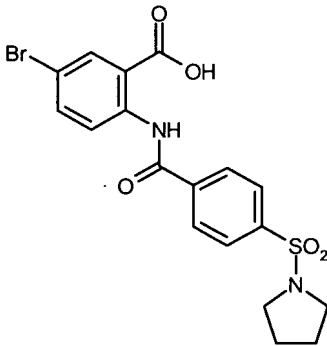
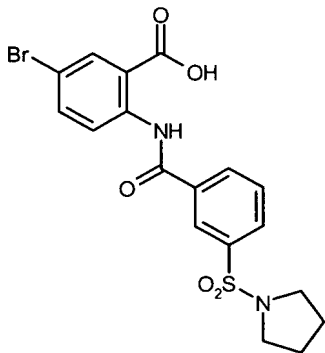
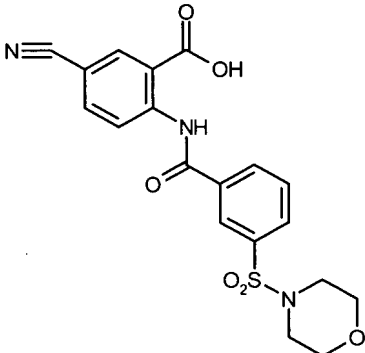
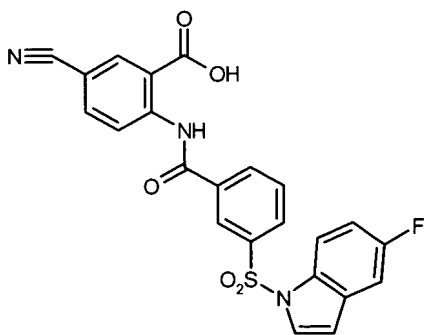
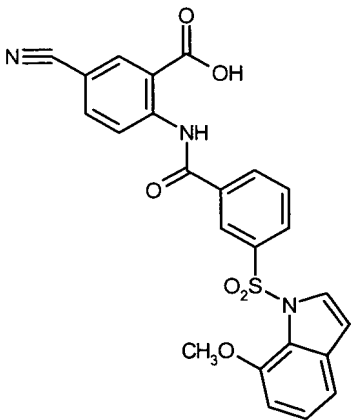
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-562862 	4	PHA-562745 	0.25
PHA-562863 	2	PHA-563275 	2
PHA-563274 	2	PHA-563277 	2

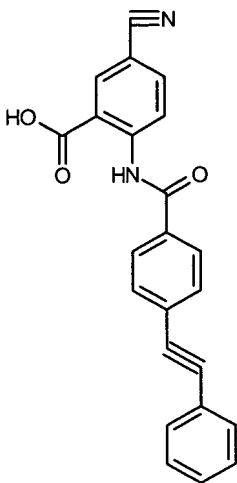
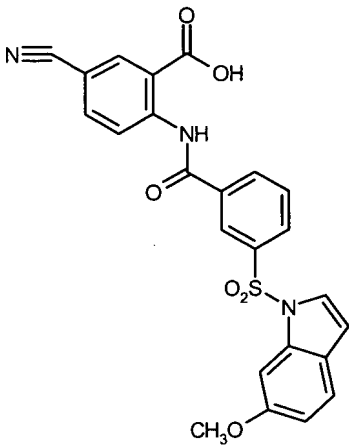
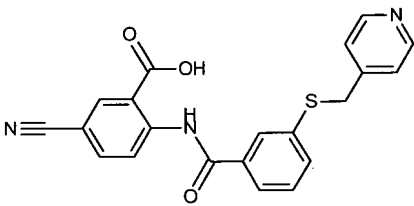
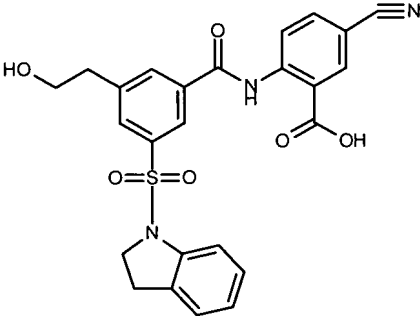
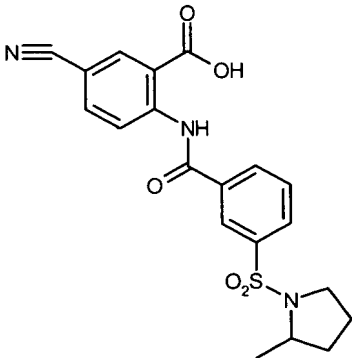
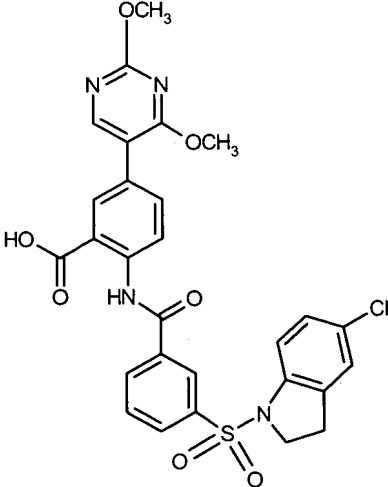


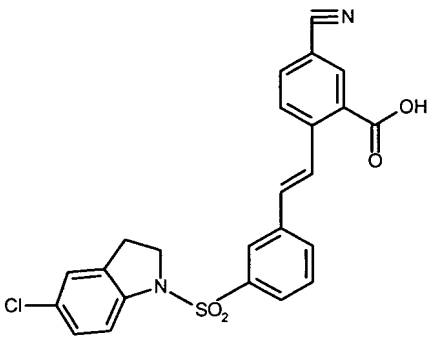
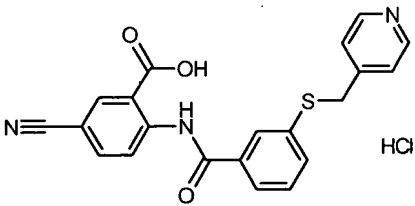
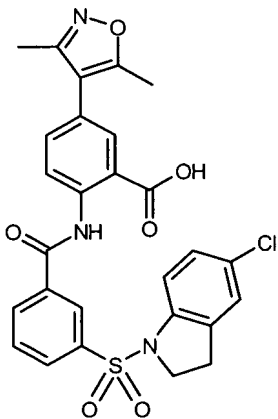
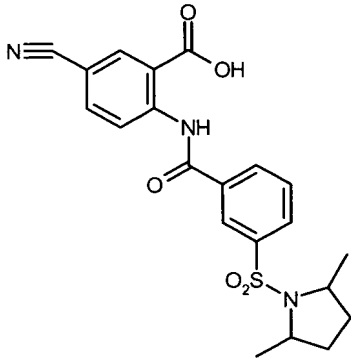
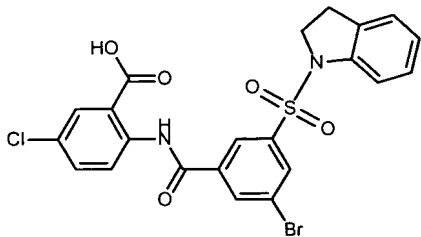
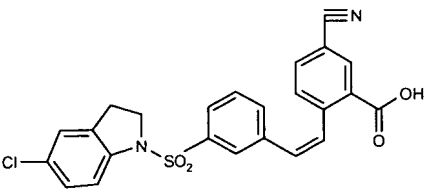
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563276 	2	PHA-563279 	0.5
PHA-563278 	2	PHA-563281 	1
PHA-563280 	1	PHA-563283 	16

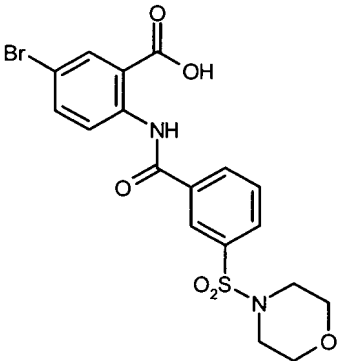
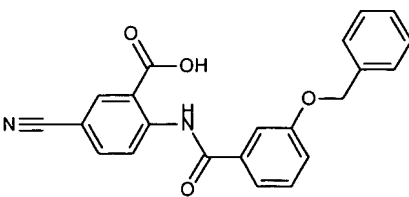
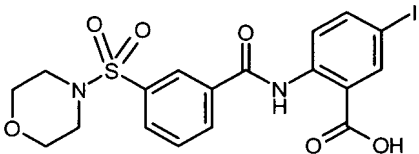
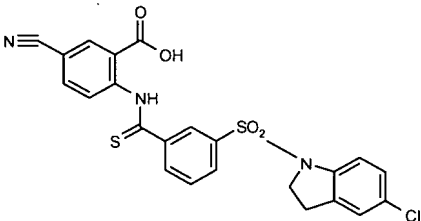
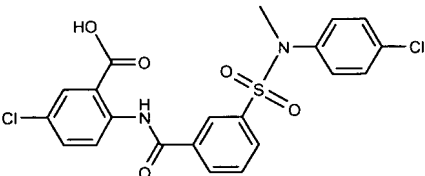
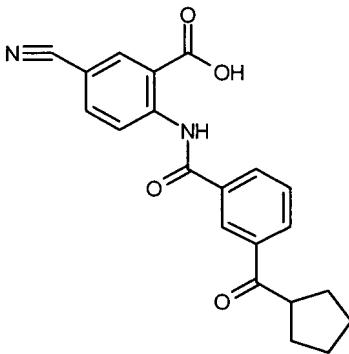
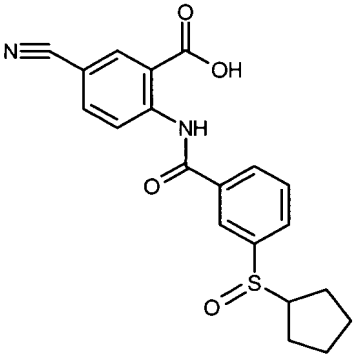
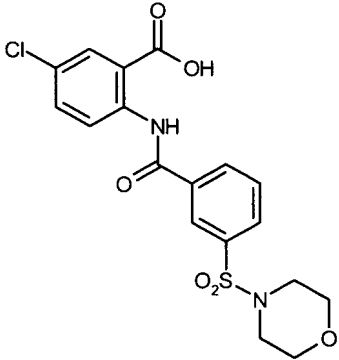
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563282 	1	PHA-563285 	2
PHA-563284 	2	PHA-564215 	0.5
PHA-563324 	>128	PHA-564750 	0.25

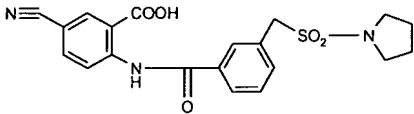
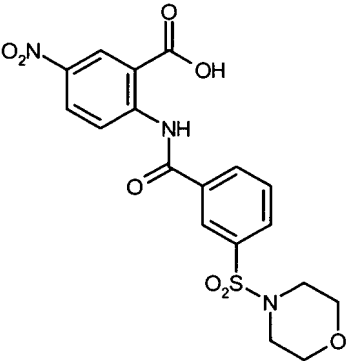
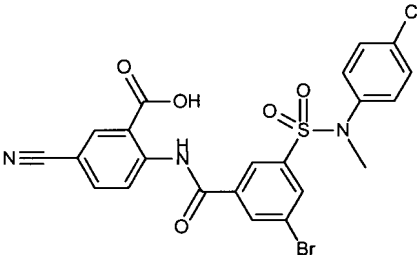
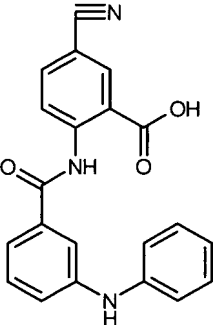
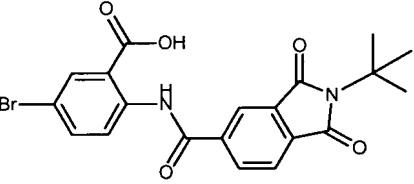
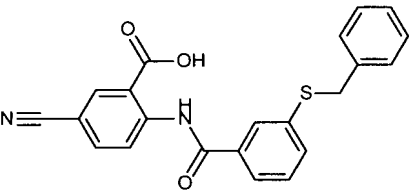
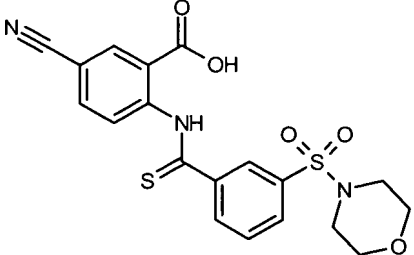
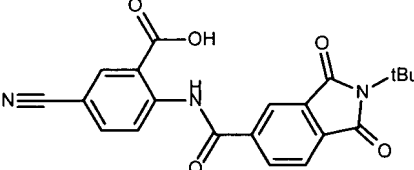
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-564218 	1	PHA-566948  (-)-enantiomer	1
PHA-566947  (+)-enantiomer	0.5	PHA-568197  6.3/93.7 trans/cis	16
PHA-568196  98/2 mixture of trans/cis	1	PHA-568205 	2

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-568206 	2	PHA-568376 	16
PHA-568378 	2	PHA-568420 	0.5
PHA-568461 	0.125	PHA-568422 	0.125

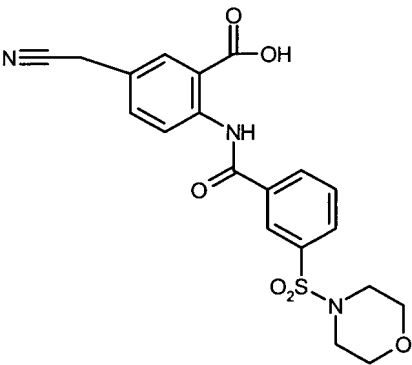
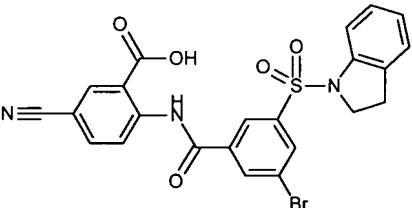
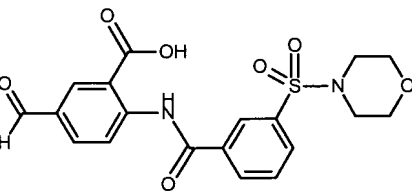
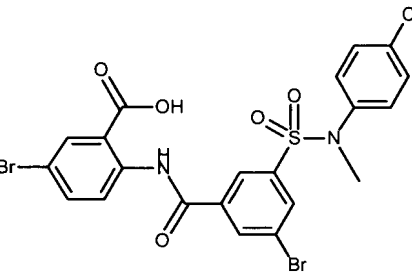
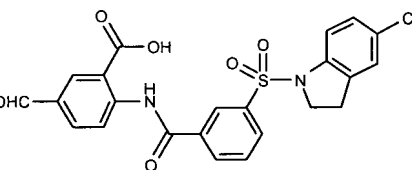
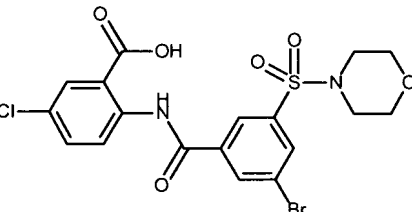
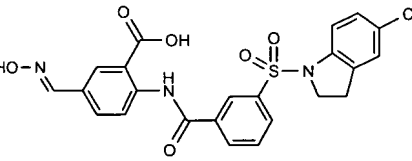
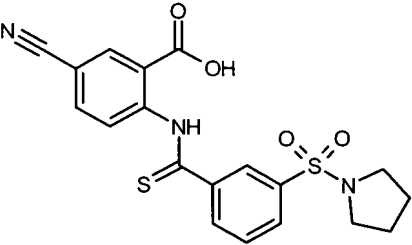
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-568907 	8	PHA-568424 	1
PHA-569044 	0.25	PHA-568425 	8
PHA-569064 	1	PHA-568906 	8

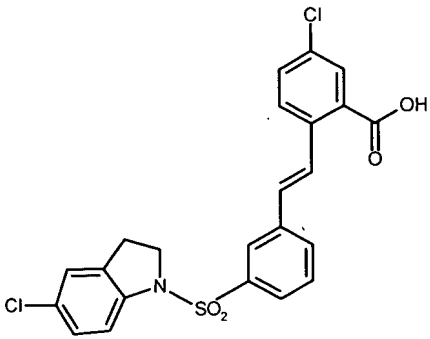
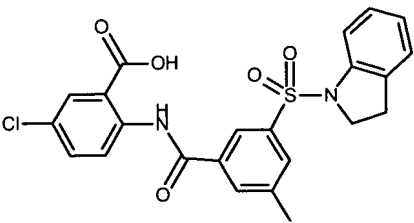
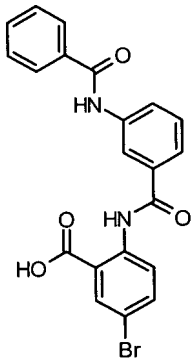
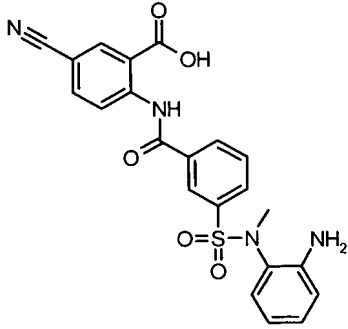
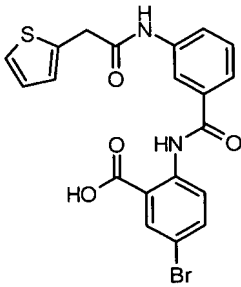
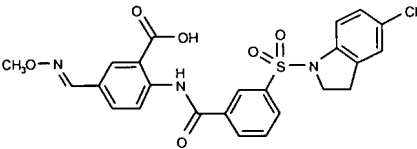
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-569887  Trans	0.25	PHA-569044A  HCl	0.5
PHA-569977 	16	PHA-569077 	1
PHA-570949 	1	PHA-569885  This is 97.9/2.1 cis/trans	16

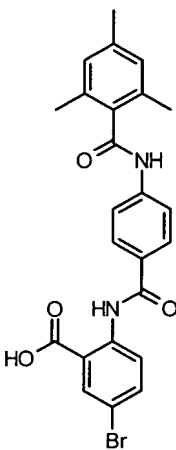
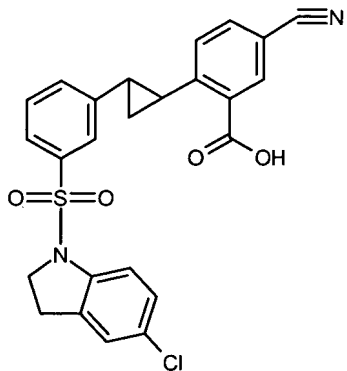
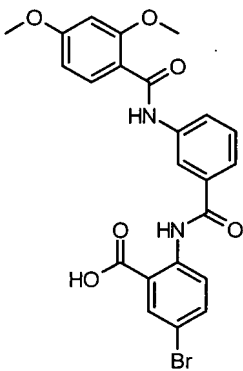
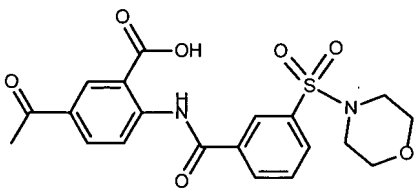
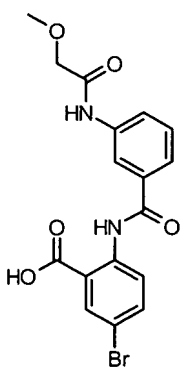
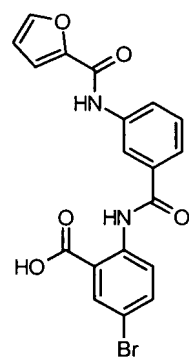
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571396 	4	PHA-569974 	1
PHA-571458 	4	PHA-570008 	0.125
PHA-615551 	1	PHA-570042 	2
PHA-630427 	4	PHA-571395 	4

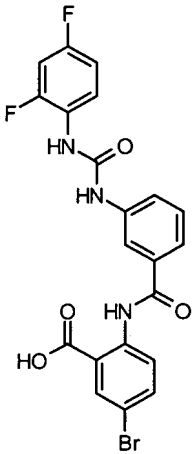
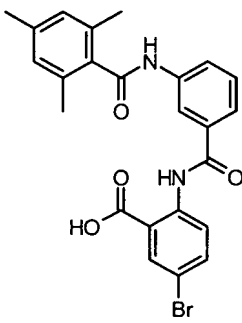
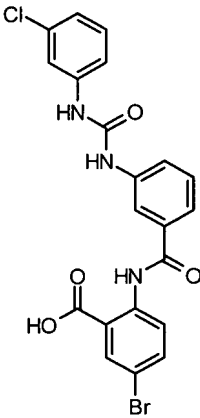
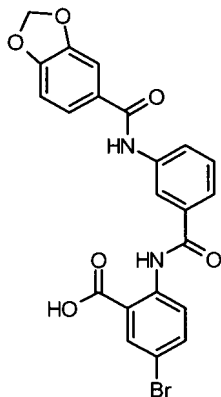
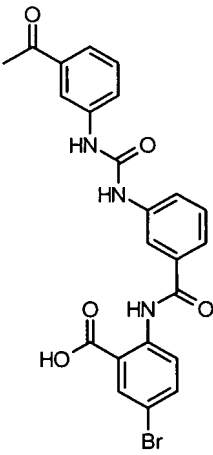
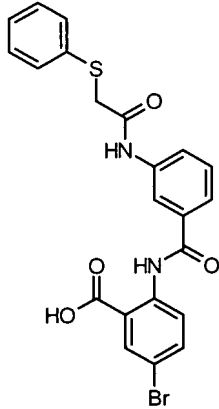
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-630852 	4	PHA-571397 	4
PHA-630966 	0.25	PHA-610938 	1
PHA-630989 	4	PHA-630368 	0.5
PHA-662430 	1	PHA-630726 	4

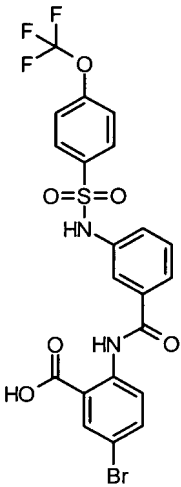
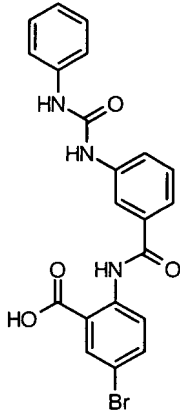
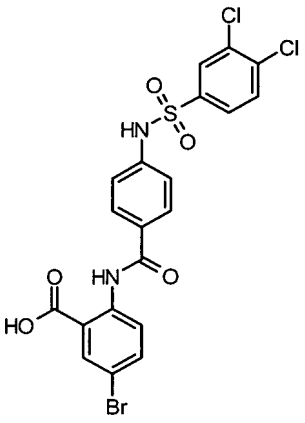
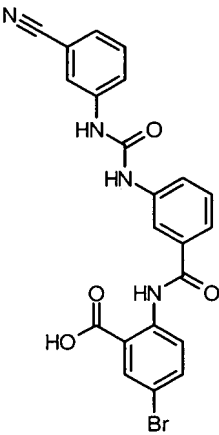
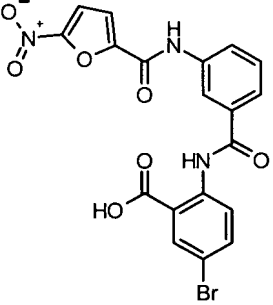
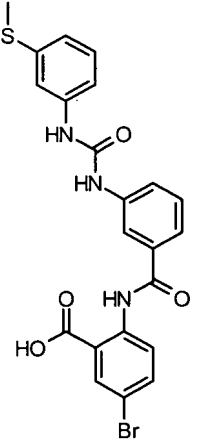


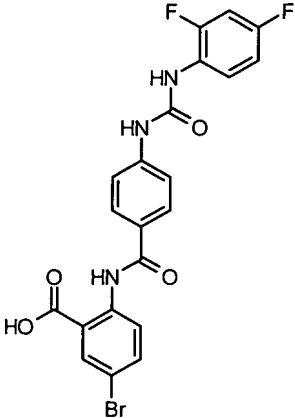
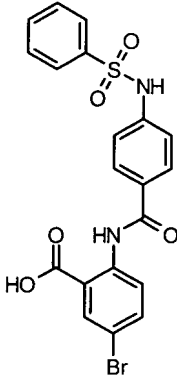
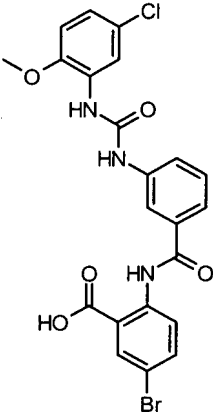
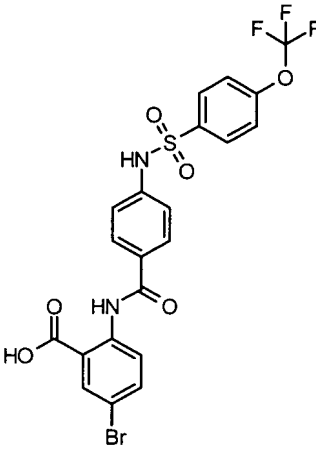
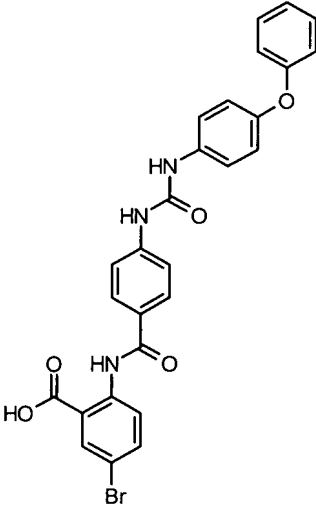
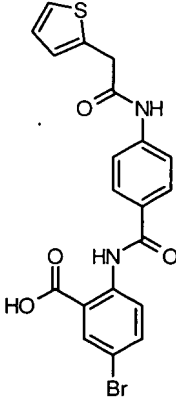
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-662951 	32	PHA-630965 	0.25
PHA-666124 	32	PHA-631082 	0.25
PHA-681768 	1	PHA-662250 	1
PHA-686834 	4	PHA-662431 	1

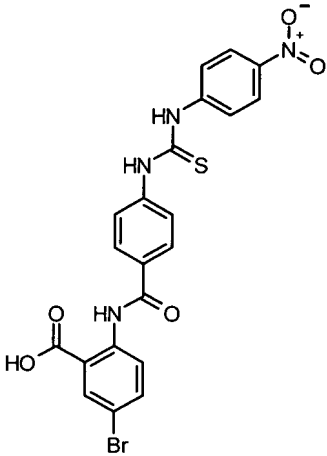
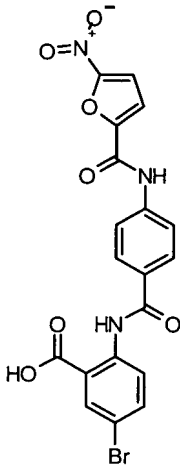
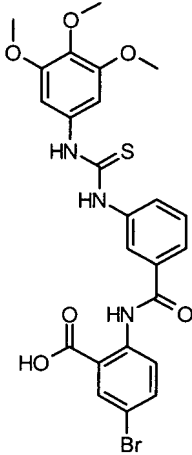
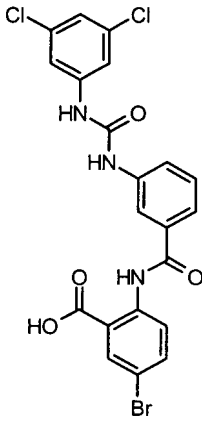
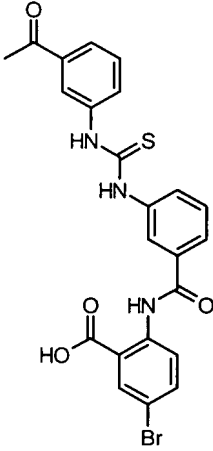
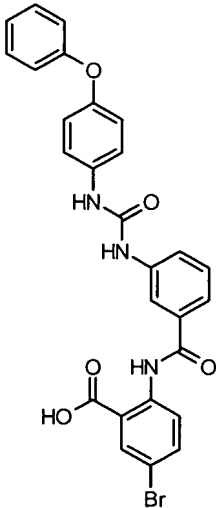
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-707801 	4	PHA-664658 	4
PHA-708976 	32	PHA-670083 	0.5
PHA-708980 	16	PHA-682996 	64

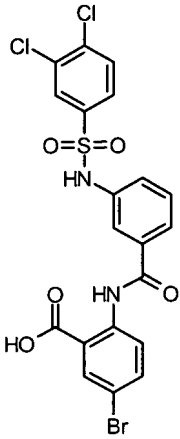
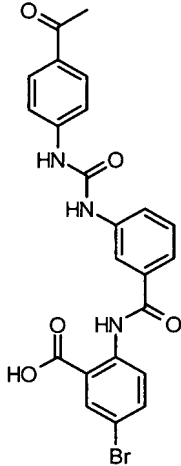
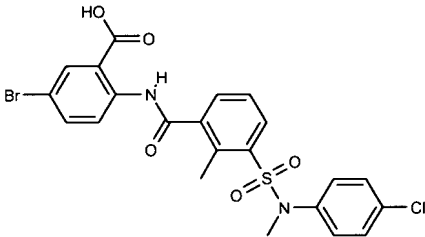
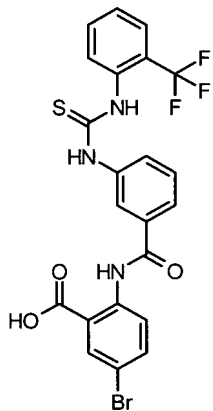
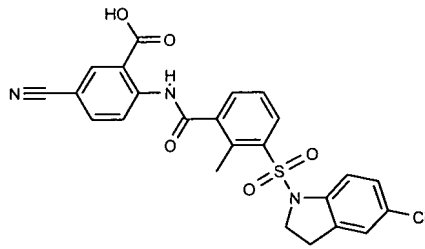
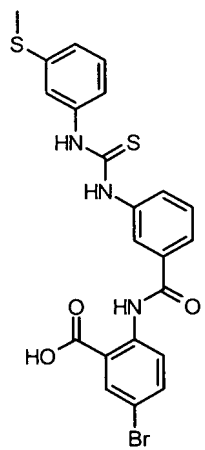
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-708982 	128	PHA-687511  is >99 trans	4
PHA-708984 	32	PHA-708923 	32
PHA-708986 	64	PHA-708978 	32

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-708989 	8	PHA-708981 	16
PHA-708991 	4	PHA-708983 	32
PHA-708993 	4	PHA-708985 	8

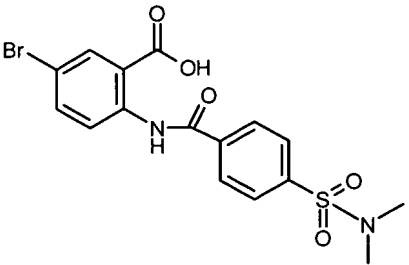
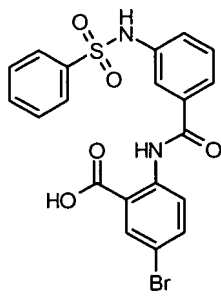
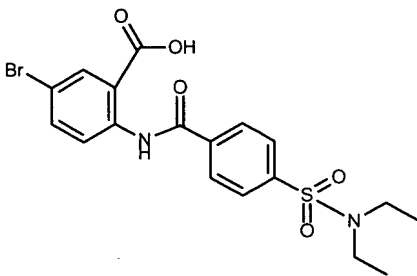
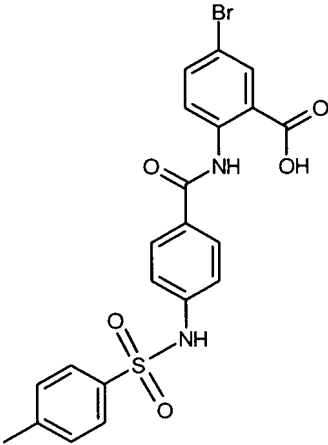
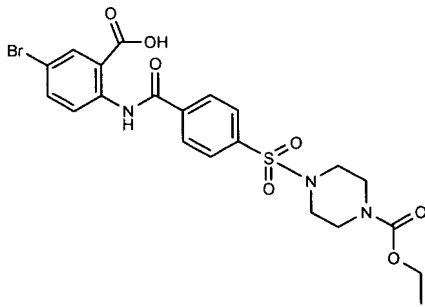
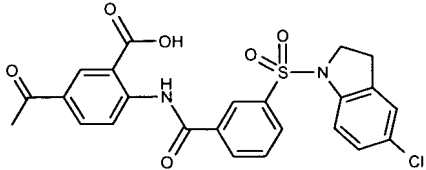
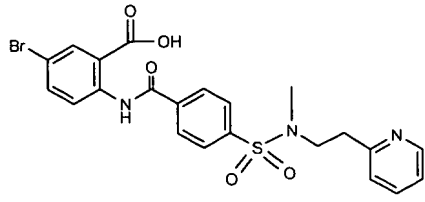
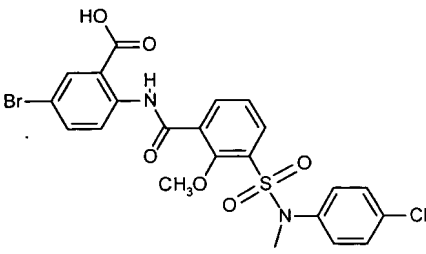
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-708995 	0.125	PHA-708988 	32
PHA-708997 	8	PHA-708990 	8
PHA-713387 	128	PHA-708992 	4

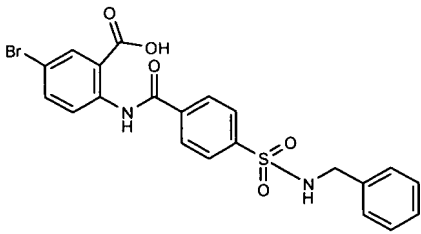
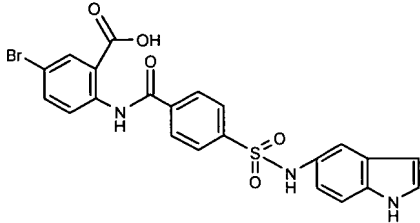
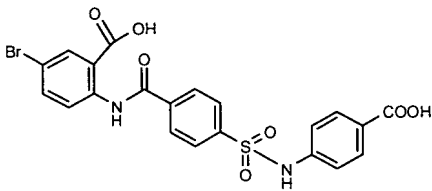
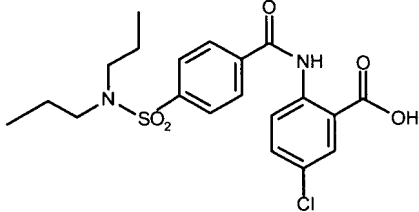
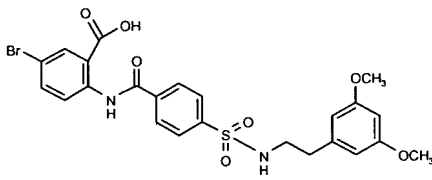
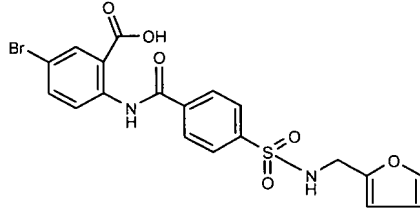
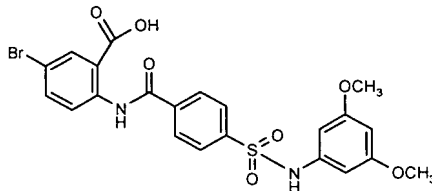
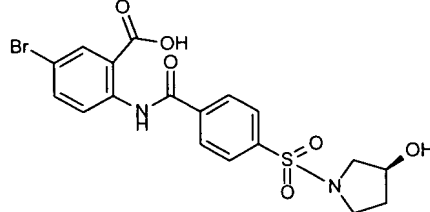
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-713394 	128	PHA-708994 	8
PHA-713398 	4	PHA-708996 	16
PHA-713400 	16	PHA-713386 	128

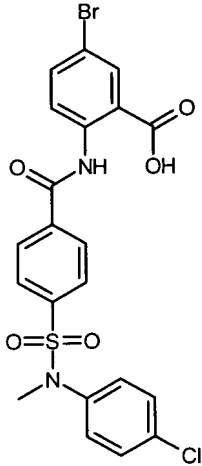
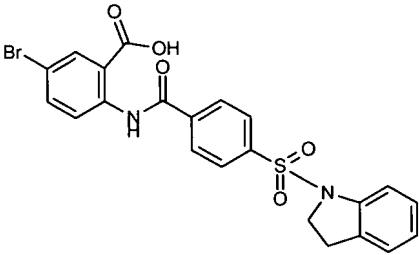
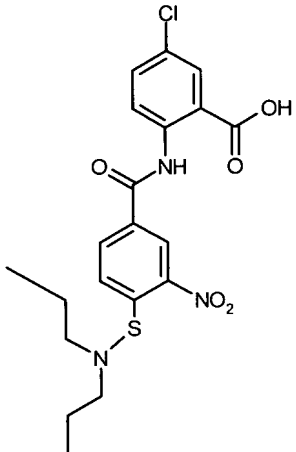
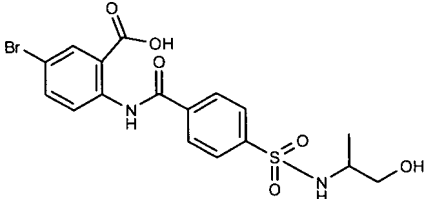
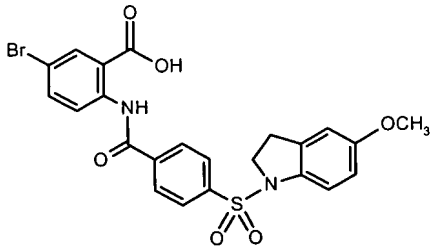
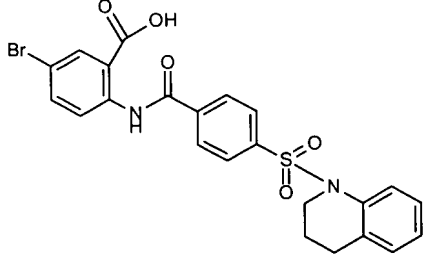
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-713403 	64	PHA-713388 	16
PHA-713406 	64	PHA-713396 	8
PHA-713408 	64	PHA-713399 	16

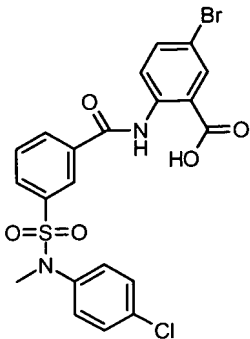
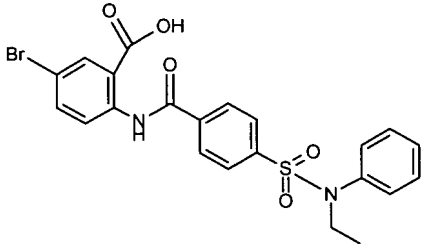
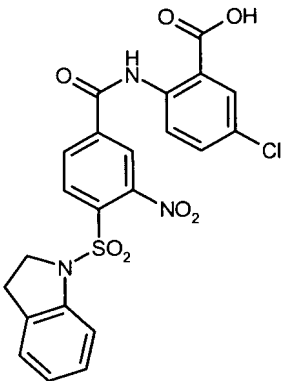
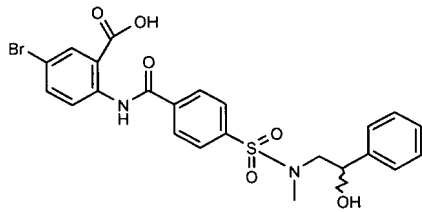
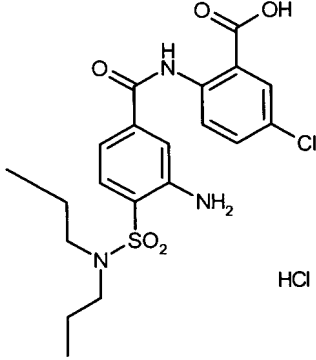
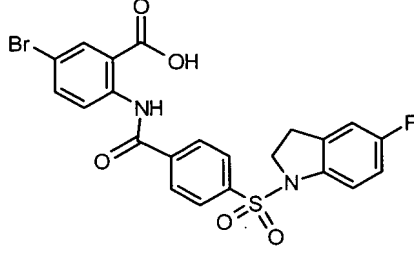
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-713410 	1	PHA-713401 	32
PHA-717196 	4	PHA-713405 	128
PHA-728844 	0.25	PHA-713407 	32

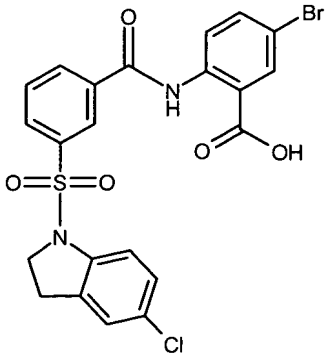
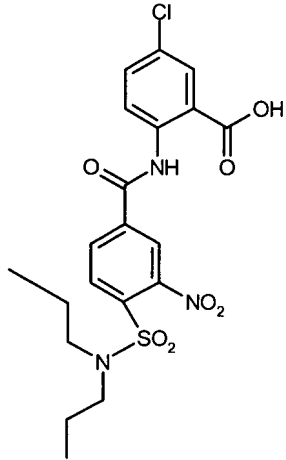
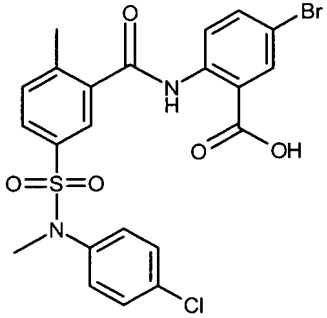
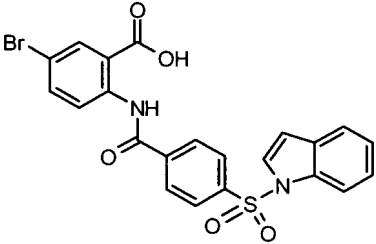
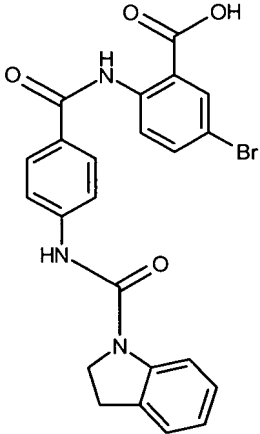
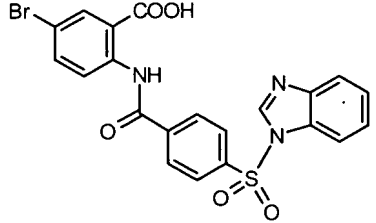


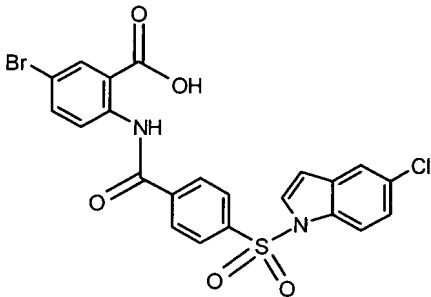
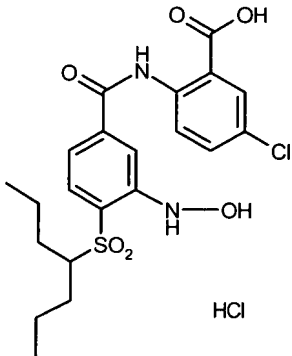
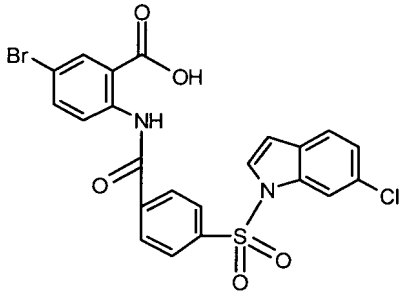
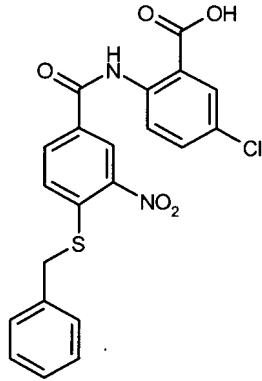
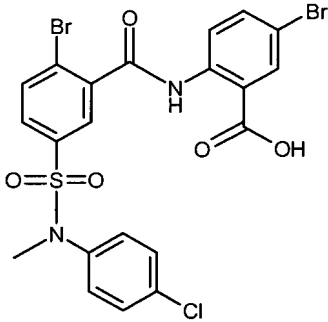
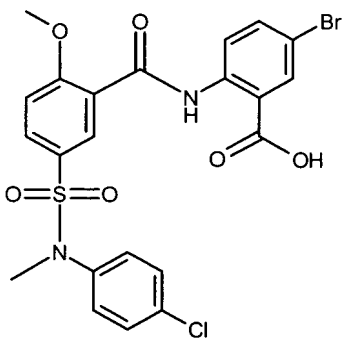
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PNU-263533 		PHA-713409 	1
PNU-271584 		PHA-713411 	32
PNU-276296 		PHA-719201 	2
PNU-276637 		PHA-735753 	16

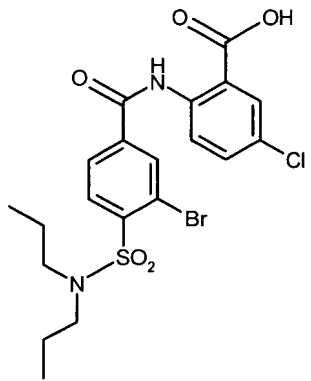
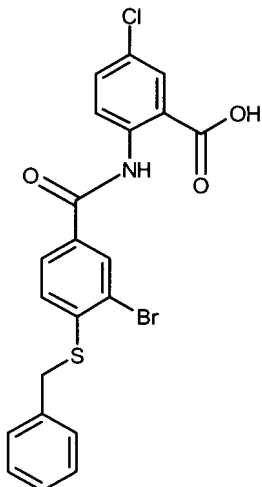
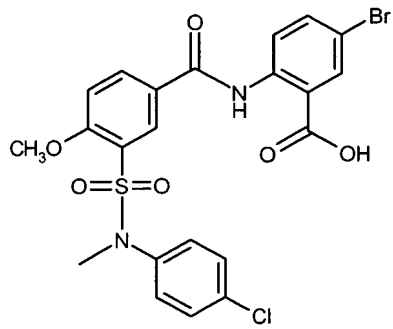
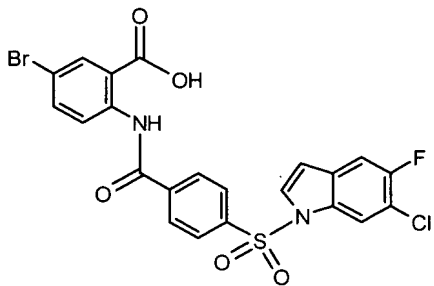
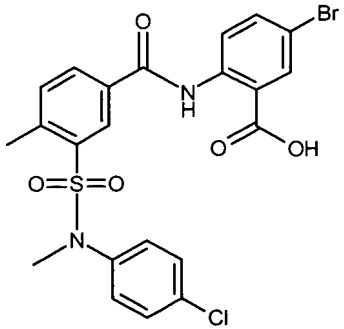
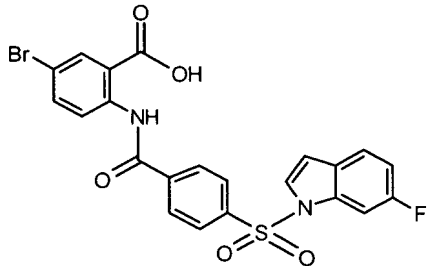
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
<p>PNU-276670</p>  <p>C<sub>21</sub>H<sub>17</sub>BrN<sub>2</sub>O<sub>5</sub>S Exact wt. 488.0042</p>		<p>PNU-268205</p> 	
<p>PNU-276817</p> 	4	<p>PNU-275747</p> 	
<p>PNU-276854</p> 		<p>PNU-276301</p> 	
<p>PNU-276933</p> 		<p>PNU-276638</p>  <p>C<sub>18</sub>H<sub>17</sub>BrN<sub>2</sub>O<sub>6</sub>S Exact wt. 467.9991</p>	

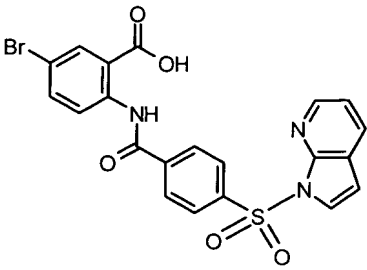
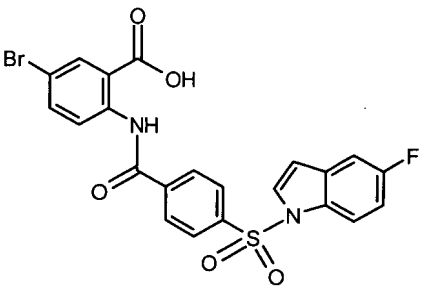
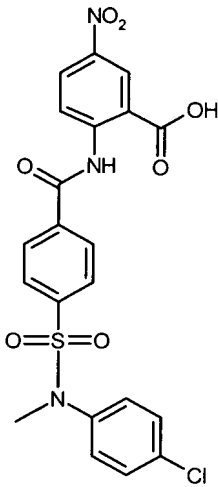
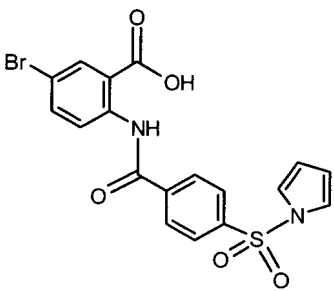
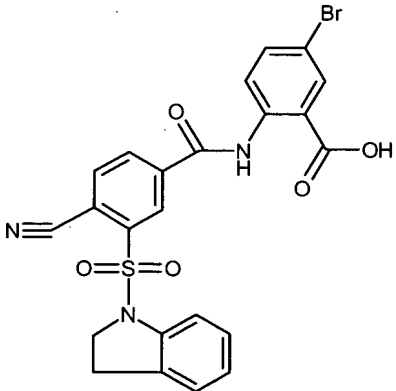
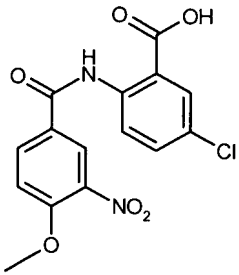
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
<p>PNU-276988</p> 	16	<p>PNU-276728</p>  <p>C<sub>22</sub>H<sub>17</sub>BrN<sub>2</sub>O<sub>5</sub>S Exact wt. 500.0042</p>	2
<p>PNU-277231</p> 	1	<p>PNU-276770</p>  <p>C<sub>17</sub>H<sub>17</sub>BrN<sub>2</sub>O<sub>6</sub>S Exact wt. 455.9991</p>	
<p>PNU-280772</p> 		<p>PNU-276818</p> 	

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PNU-283076 	1	PNU-276913 	
PNU-283599 	1	PNU-276952  racemic	
PNU-283603A  HCl	16	PNU-280727 	

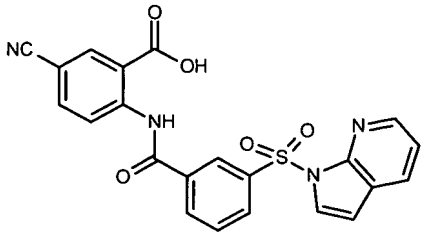
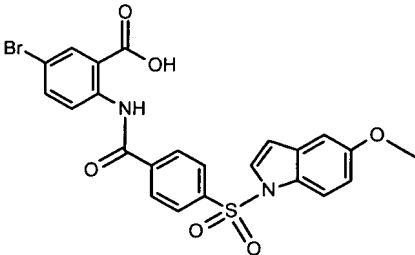
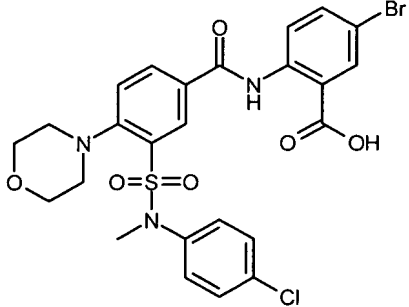
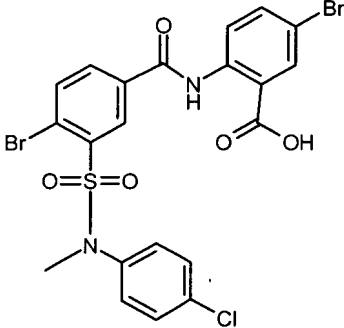
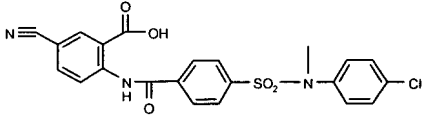
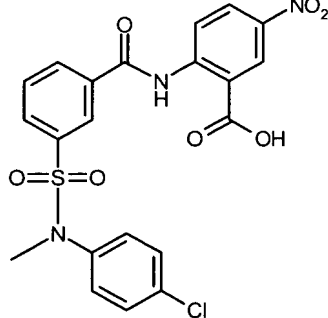
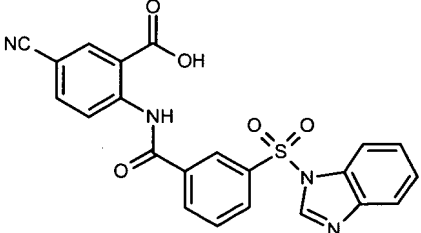
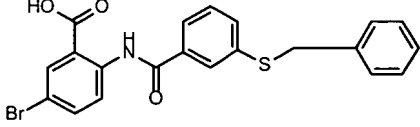
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PNU-288969 	0.25	PNU-282958 	
PNU-290821 	64	PNU-283318 	0.125
PNU-290877  <p>See Comments</p>	>128	PNU-283371 	4

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PNU-290905 	1	PNU-283601A 	32
PNU-290906 	1	PNU-283604 	4
PNU-291061 	16	PNU-289815 	8

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PNU-291410 	4	PNU-290882 	1
PNU-291570 	8	PNU-291010 	1
PNU-291571 	0.5	PNU-291011 	0.25

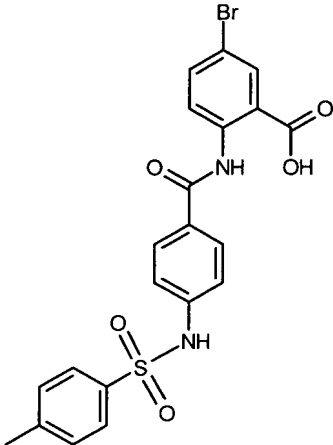
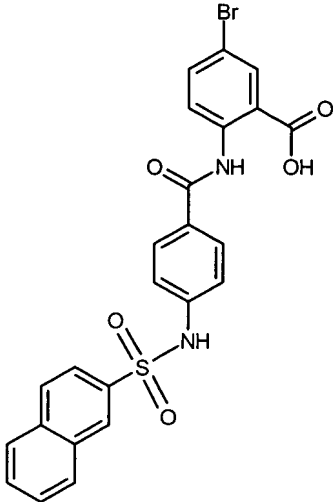
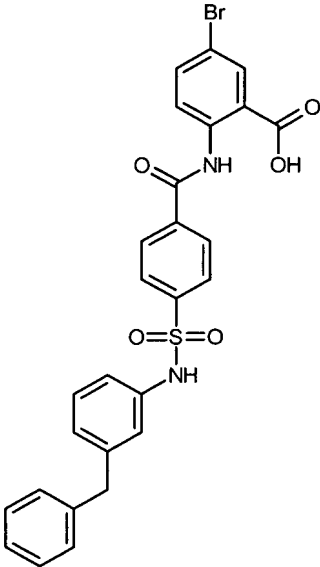
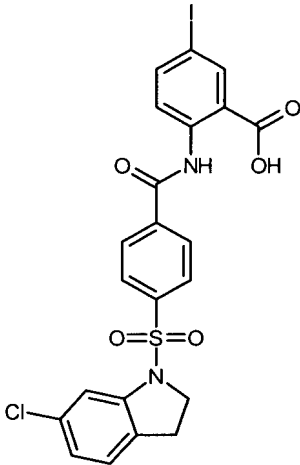
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PNU-292070 	2	PNU-291129 	0.5
PNU-293032 	16	PNU-291130 	4
PNU-293905 	8	PNU-291408 	32

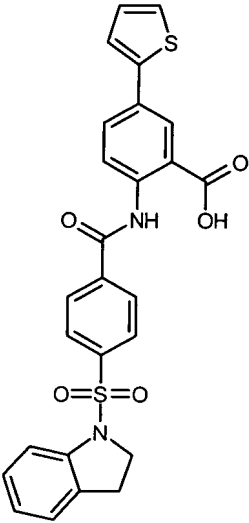
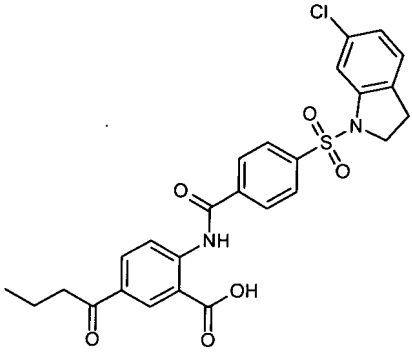
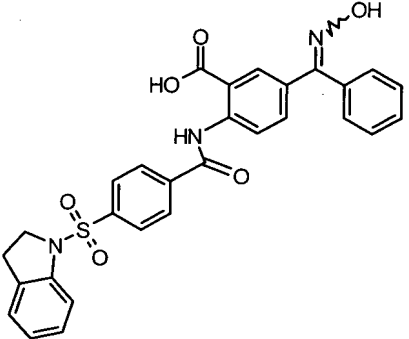
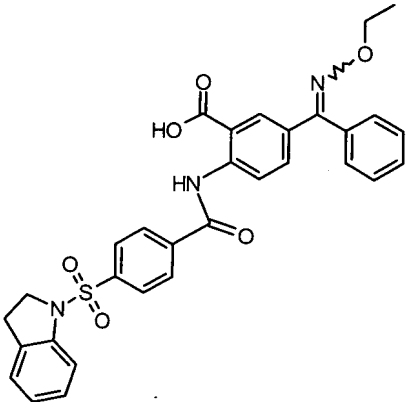
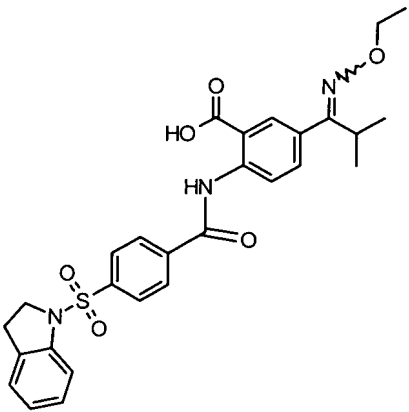
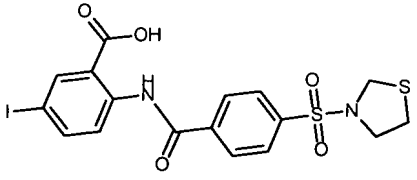


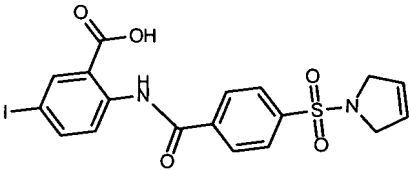
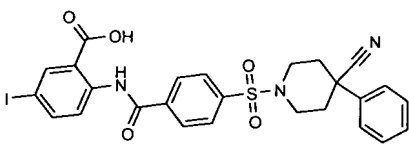
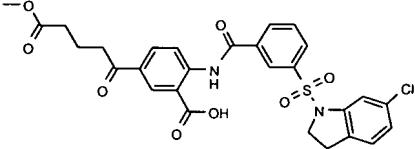
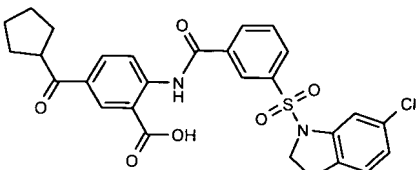
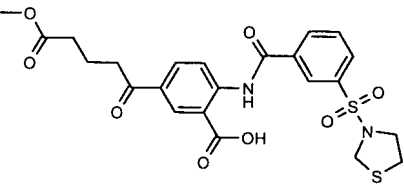
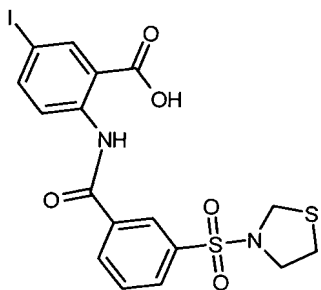
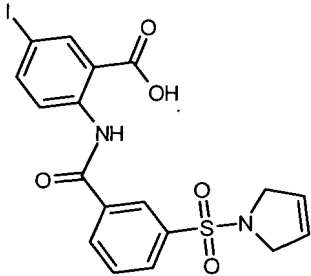
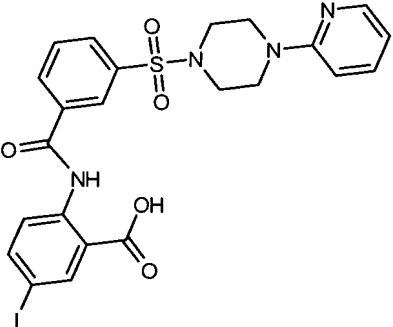
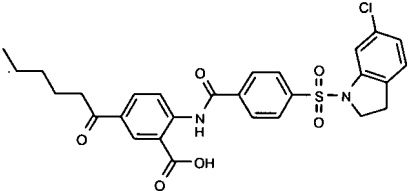
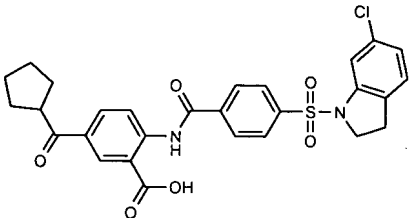
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-630331 	2	PNU-291517 	2
PNU-293795 	32	PNU-291679 	1
PNU-294595 	16	PNU-292379 	0.5
PHA-630330 	0.5	PNU-293049 	4

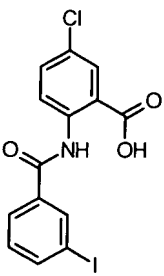
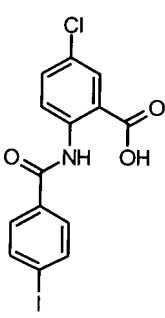
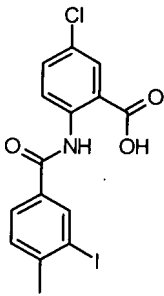
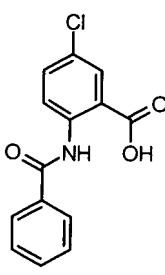
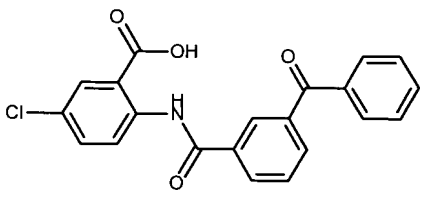
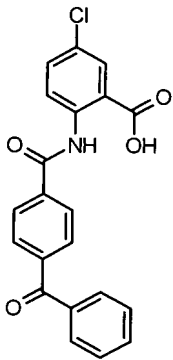
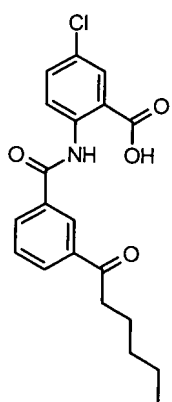
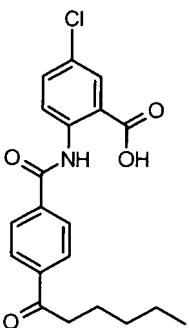
**Table 2: Activity Data**

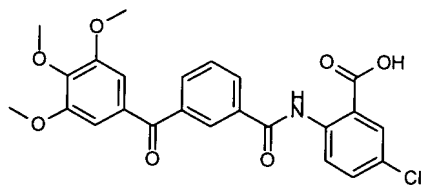
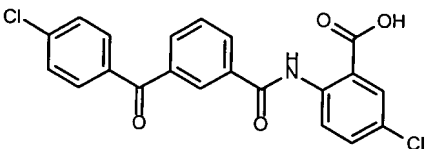
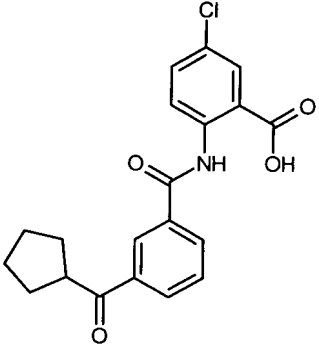
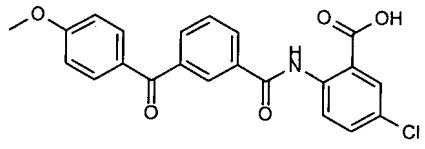
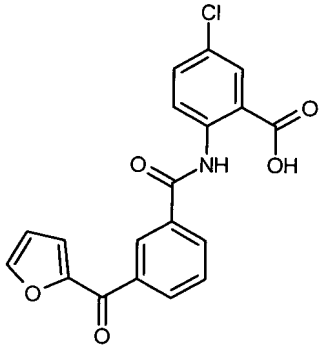
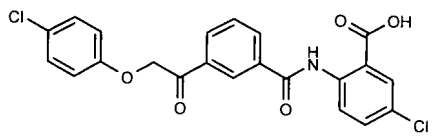
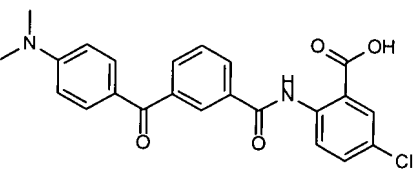
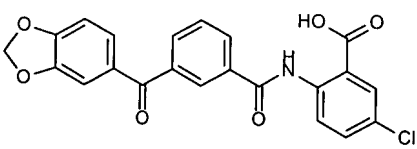
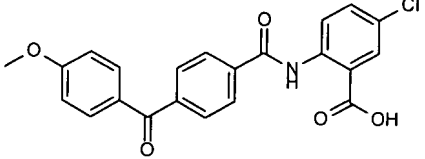
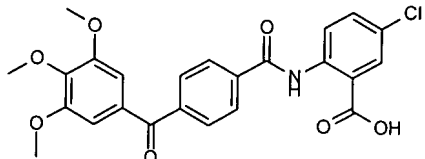
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
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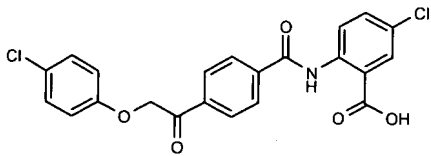
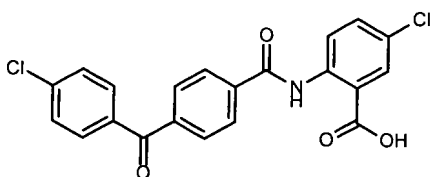
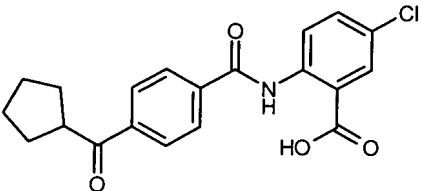
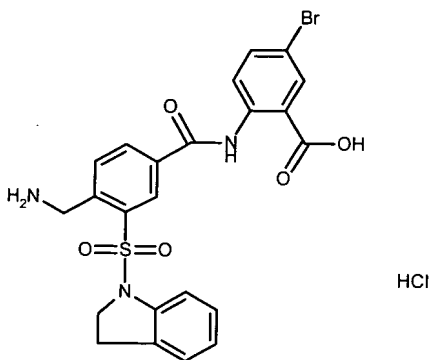
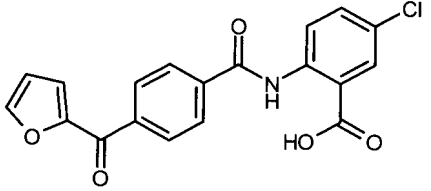
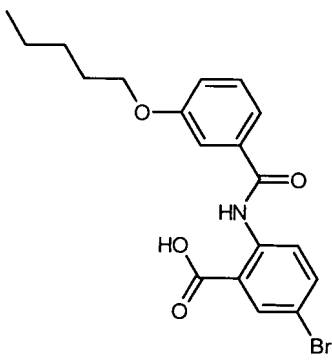
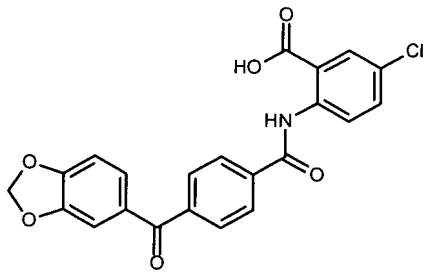
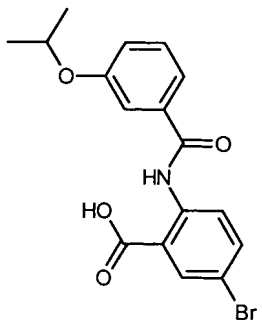
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
L-170210 	16	L-170216 	
L-199199 		L-217790 	4

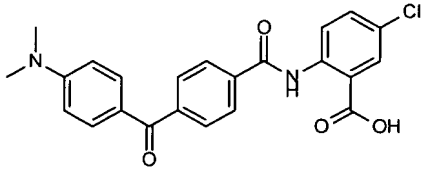
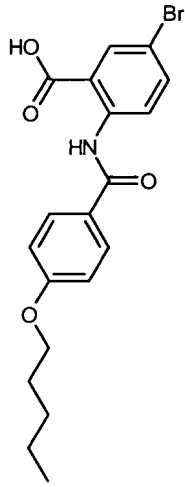
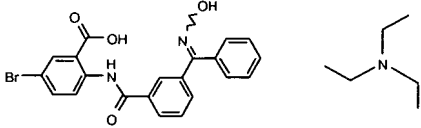
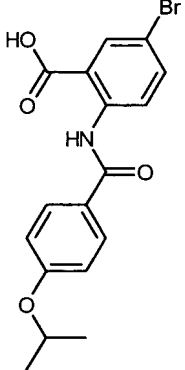
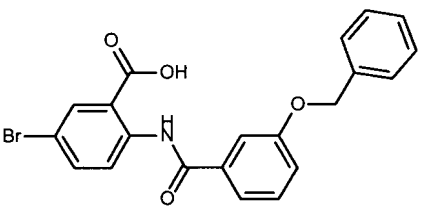
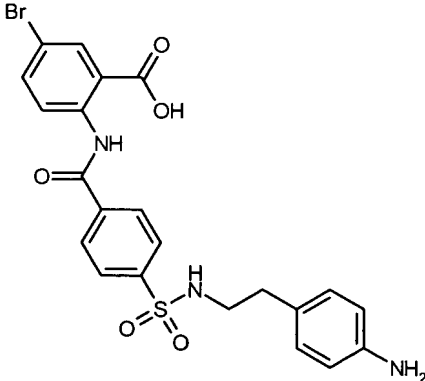
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
L-217791 	4	L-218343 	16
L-502902 	128	L-502903 	16
L-502904 	64	PHA-500140 	32

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-500152 	32	PHA-500200 	4
PHA-500218 	64	PHA-500219 	32
PHA-500230 	>128	PHA-500236 	8
PHA-500248 	8	PHA-500284 	32
PHA-502605 	8	PHA-502606 	16

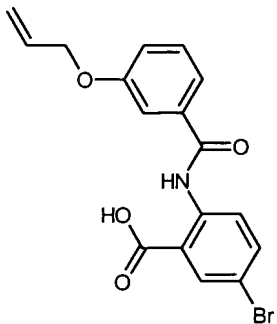
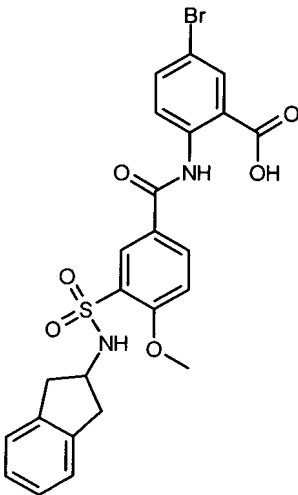
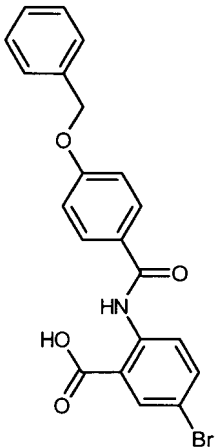
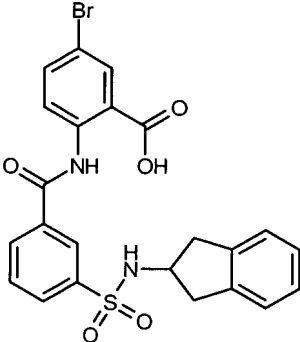
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-520185 	8	PHA-520200 	2
PHA-520221 	2	PHA-520245 	128
PHA-520412 	4	PHA-520413 	4
PHA-520414 	8	PHA-520416 	4

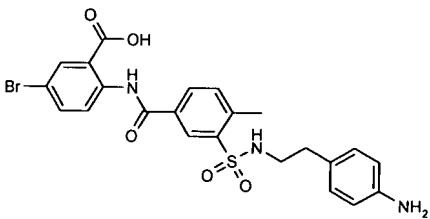
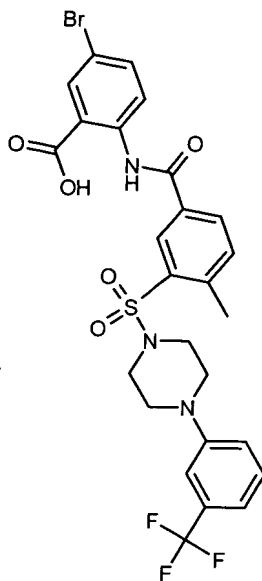
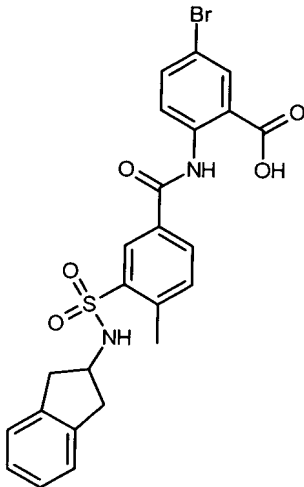
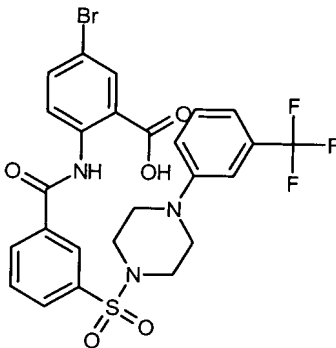
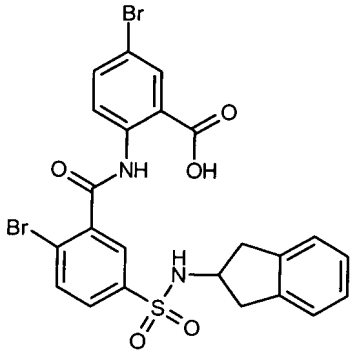
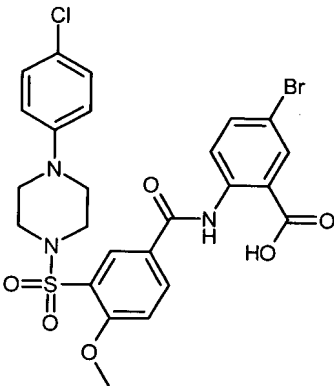
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-523506 	32	PHA-523507 	4
PHA-523510 	8	PHA-523508 	8
PHA-523511 	8	PHA-523509 	8
PHA-523513 	4	PHA-523512 	4
PHA-523516 	2	PHA-523514 	4

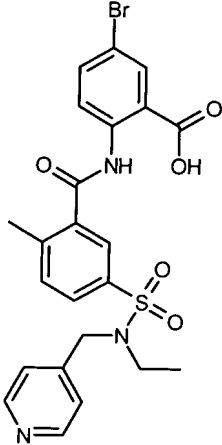
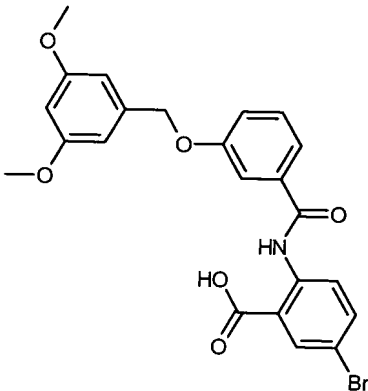
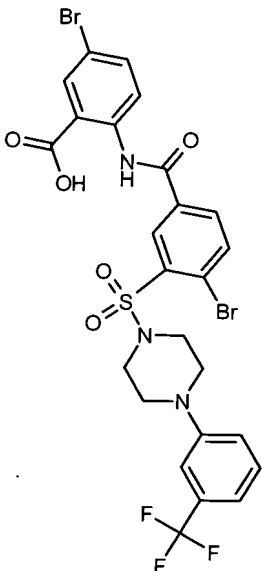
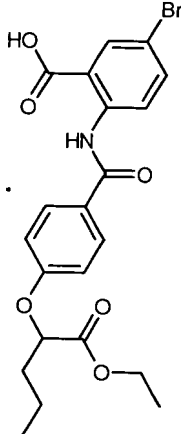
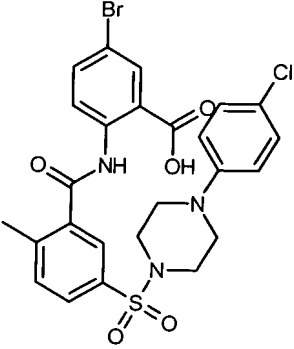
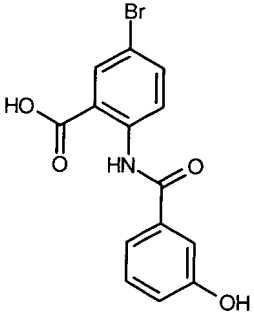
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-523517 	4	PHA-523515 	4
PHA-523518 	8	PHA-524553A  HCl	>128
PHA-523519 	4	PHA-525501 	8
PHA-523520 	2	PHA-525503 	2

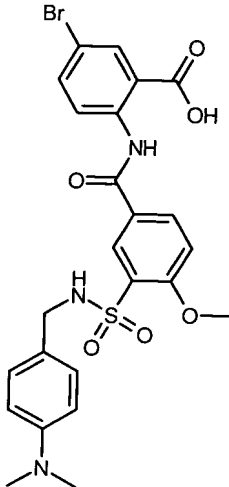
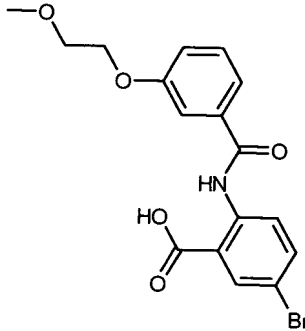
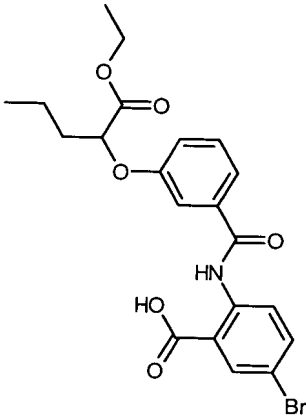
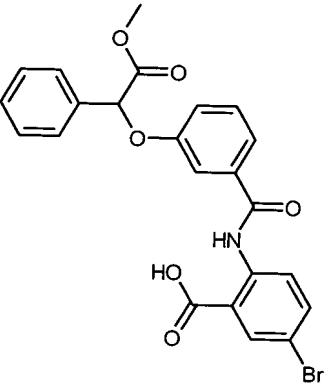
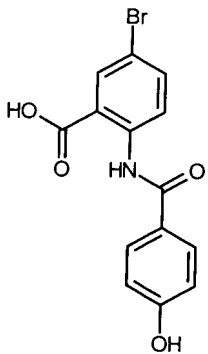
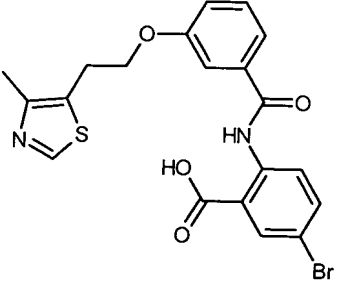
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-523521 	16	PHA-525505 	16
PHA-524545E 	0.5	PHA-525506 	64
PHA-525500 	4	PHA-526643 	64

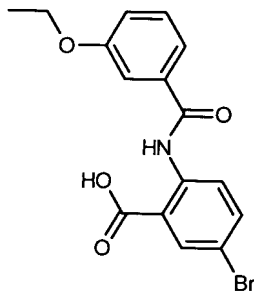
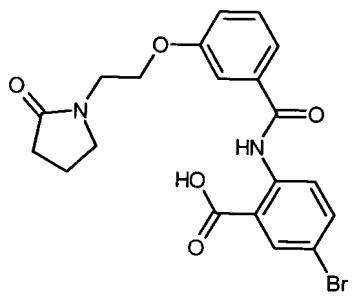
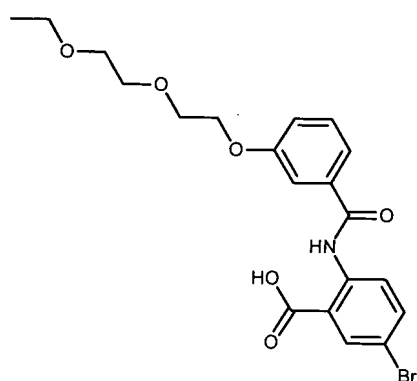
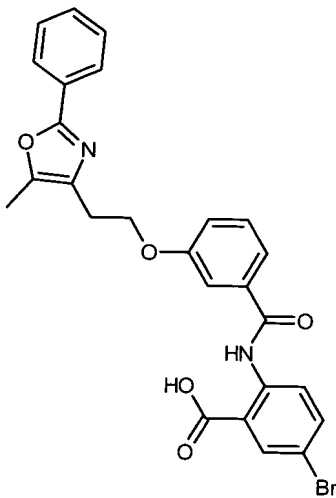
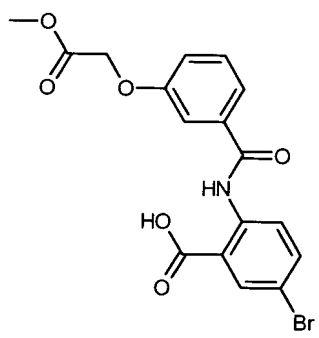
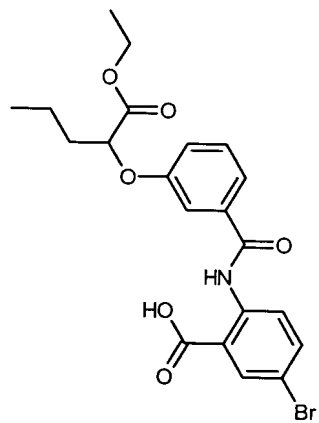


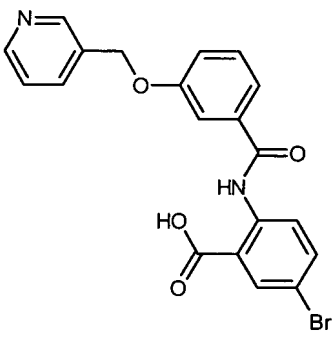
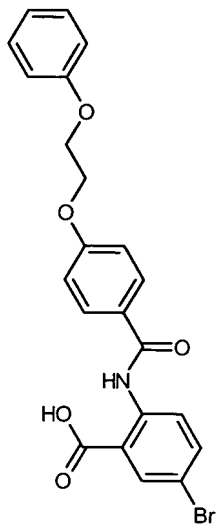
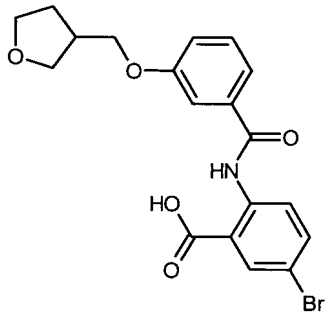
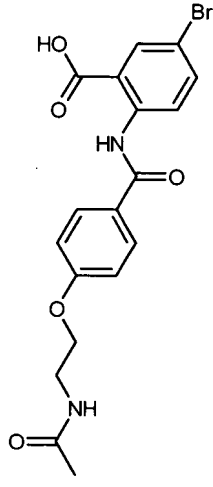
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-525502 	8	PHA-526650 	2
PHA-525504 	16	PHA-526652 	1

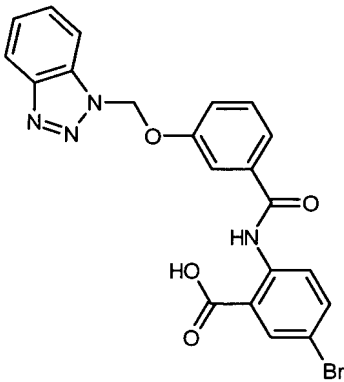
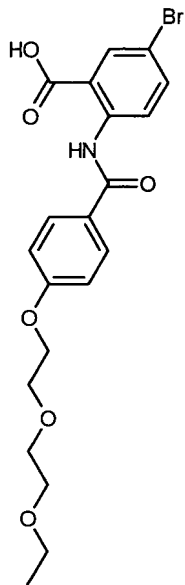
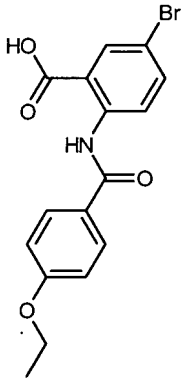
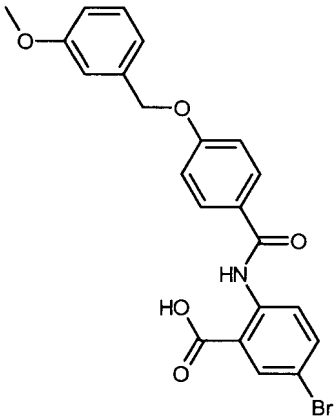
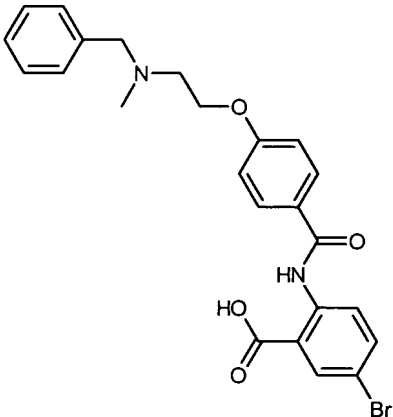
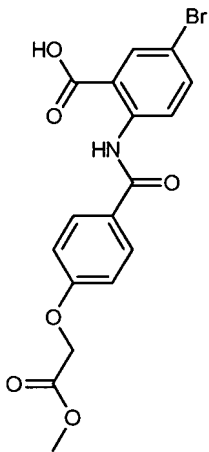
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-526641 	8	PHA-526655 	16
PHA-526648 	0.25	PHA-526661 	16
PHA-526651 	0.25	PHA-526681 	8

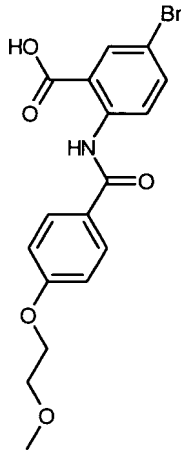
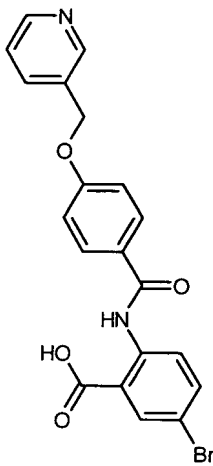
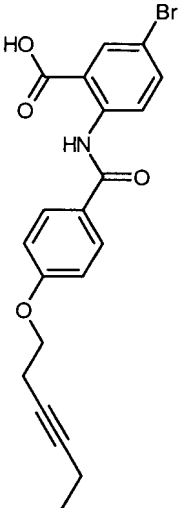
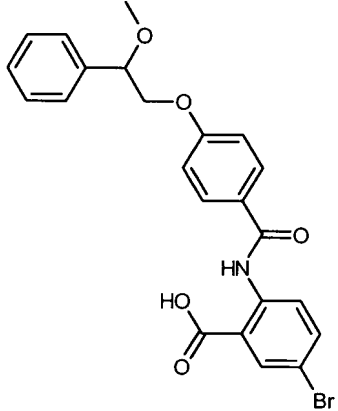
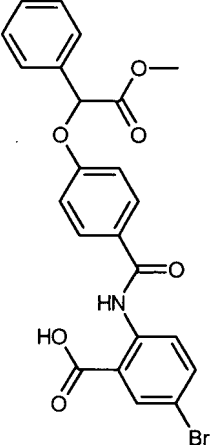
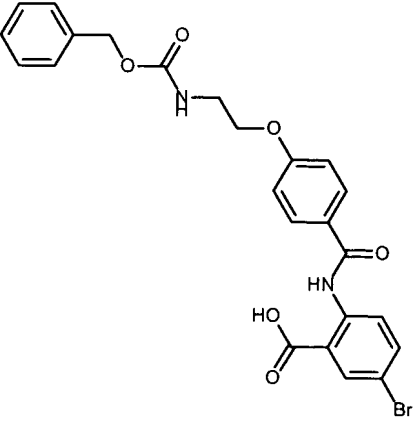
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-526653 	>128	PHA-526705 	16
PHA-526660 	8	PHA-526712 	64
PHA-526679 	32	PHA-530915 	32

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-526683 	>128	PHA-533237 	16
PHA-526707 	2	PHA-533244 	4
PHA-530914 	32	PHA-533249 	8

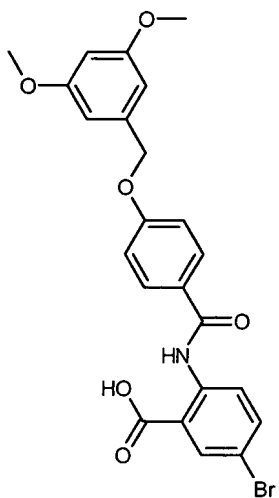
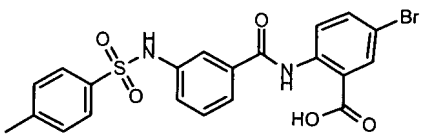
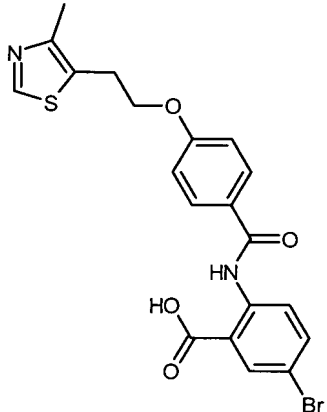
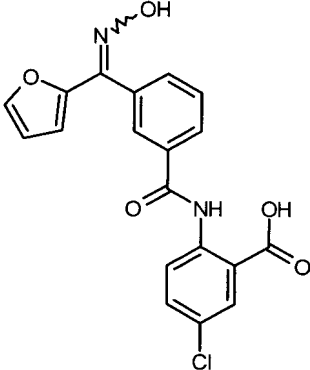
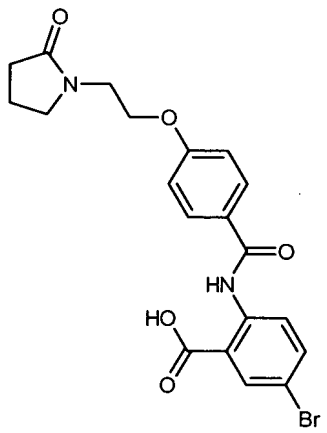
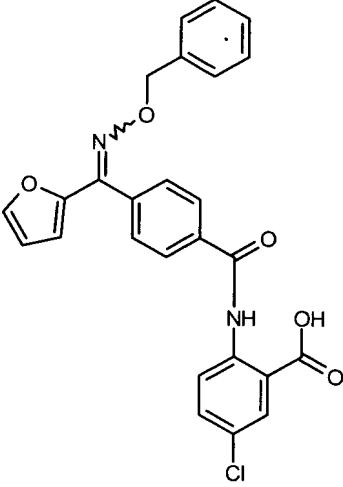
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-533232 	64	PHA-533253 	32
PHA-533243 	32	PHA-533258 	32
PHA-533247 	32	PHA-533261 	8

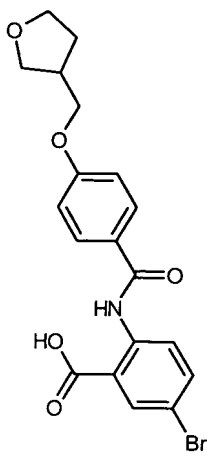
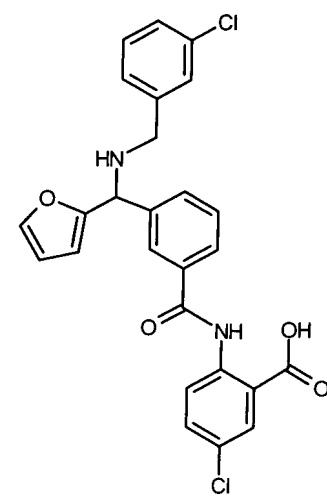
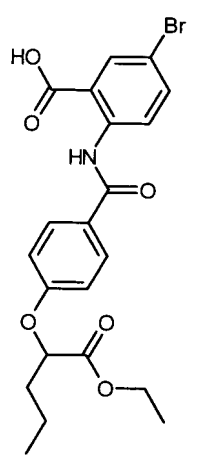
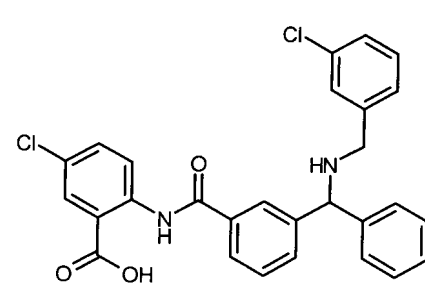
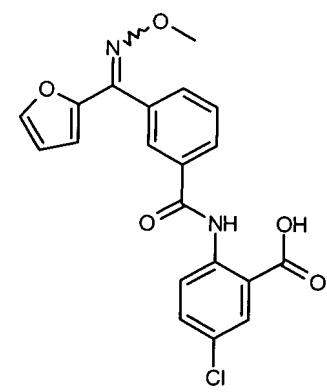
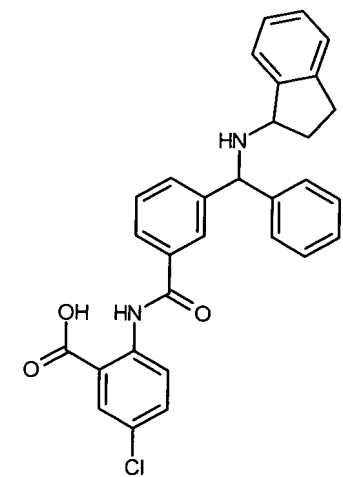
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-533252 	32	PHA-533265 	128
PHA-533257 	16	PHA-533268 	64

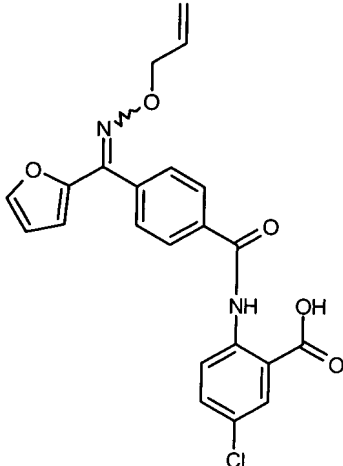
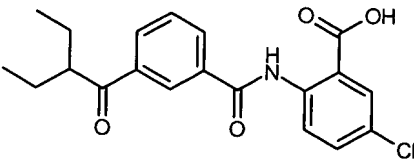
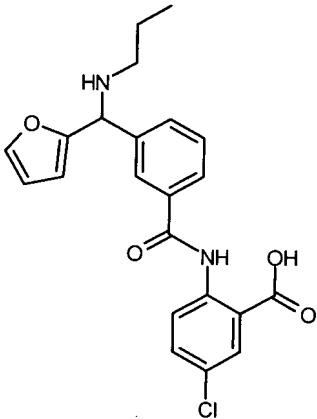
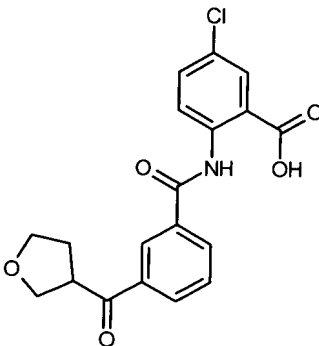
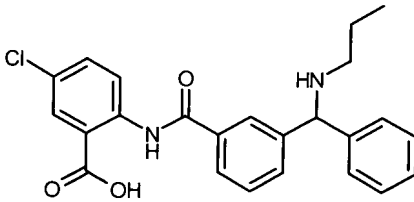
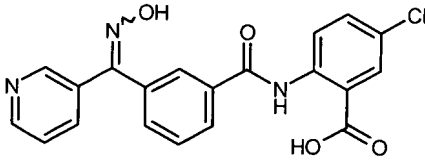
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
<p>PHA-533259</p> 	32	<p>PHA-533272</p> 	128
<p>PHA-533262</p> 	64	<p>PHA-533274</p> 	8
<p>PHA-533264</p> 	128	<p>PHA-533276</p> 	64

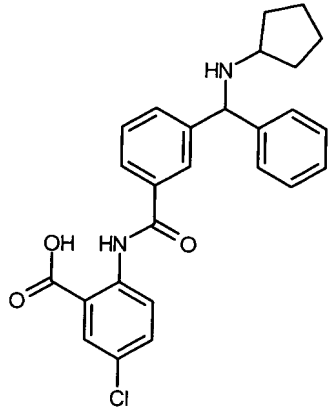
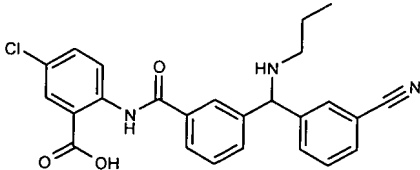
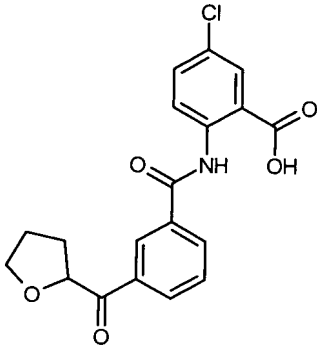
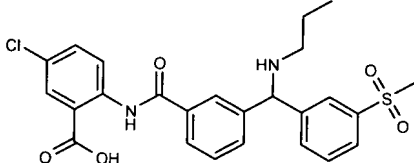
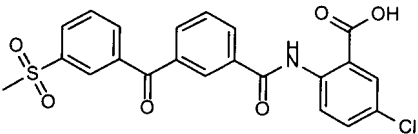
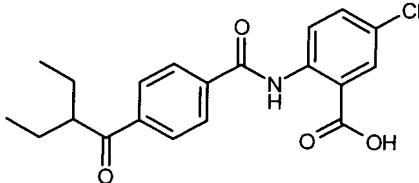
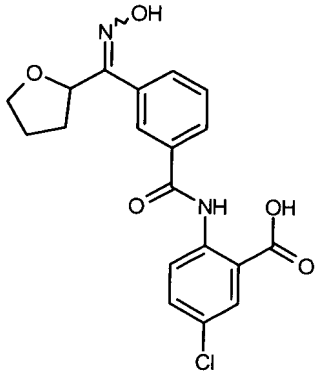
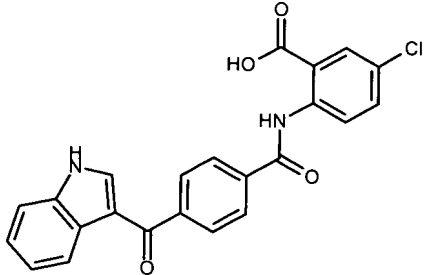
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-533266 	128	PHA-533281 	64
PHA-533269 	16	PHA-533285 	64
PHA-533273 	64	PHA-533289 	64

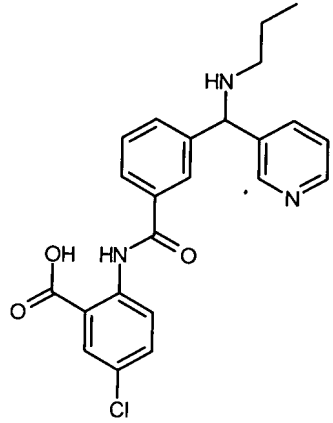
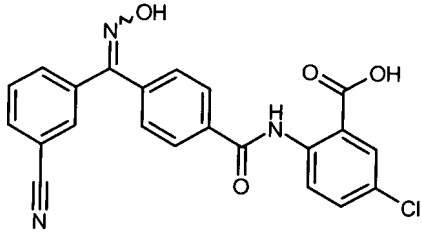
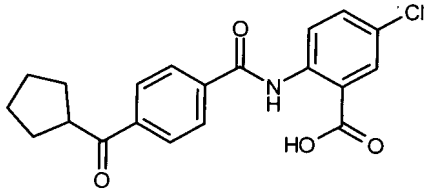
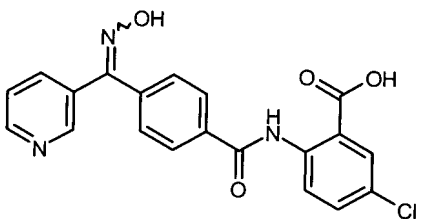
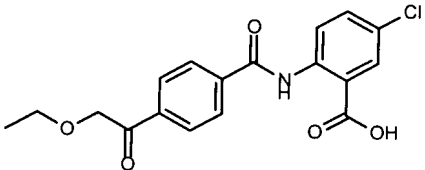
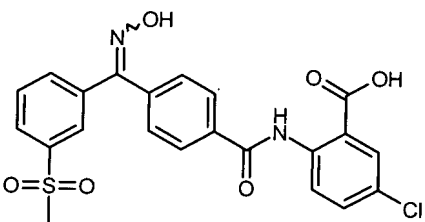
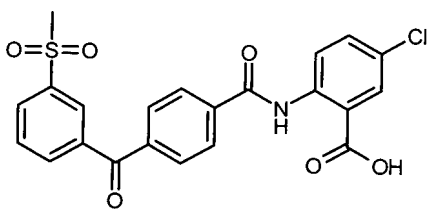
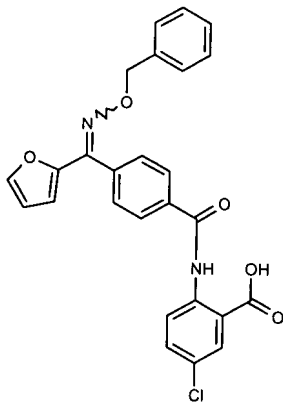


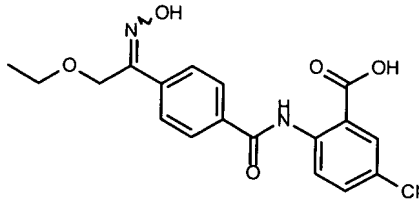
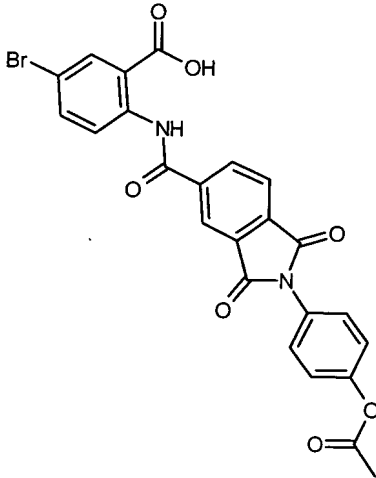
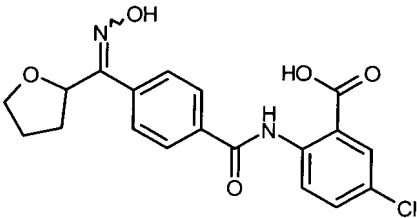
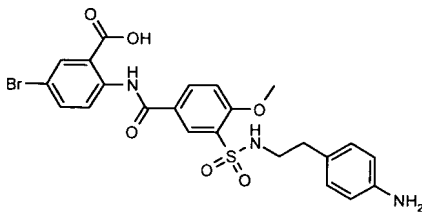
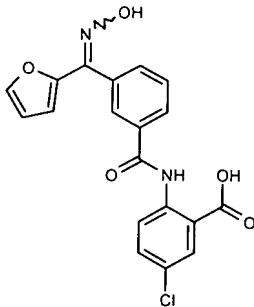
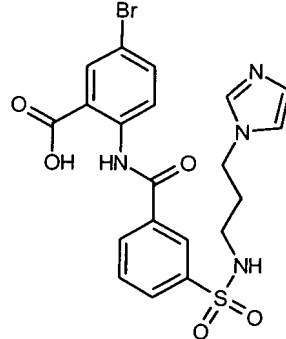
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-533275 	8	PHA-533401 	0.5
PHA-533278 	32	PHA-537084  least retained isomer by RP-HPLC	2
PHA-533282 	>128	PHA-537089  least retained isomer by RP-LC/MS	32

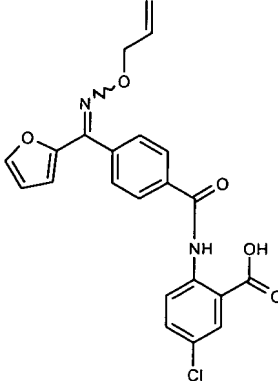
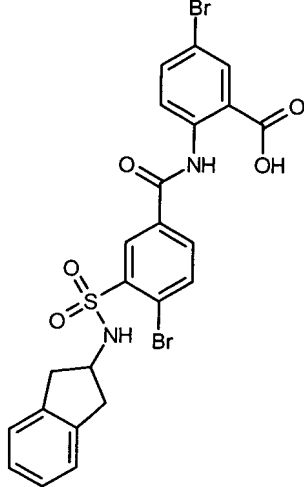
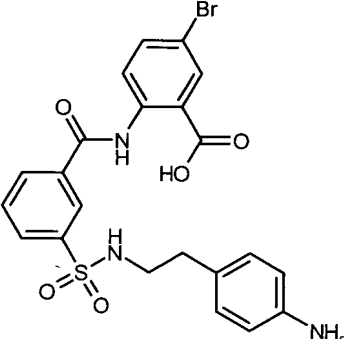
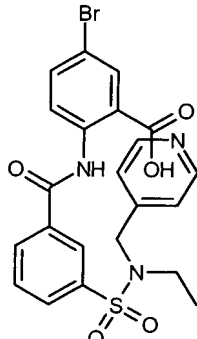
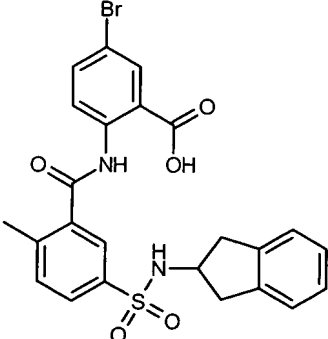
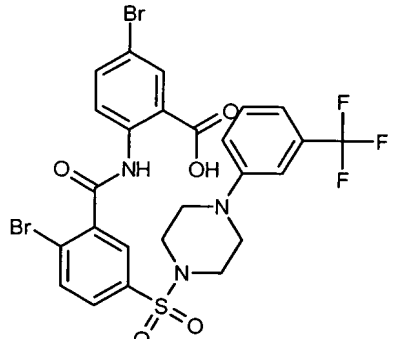
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-533286 	128	PHA-537091 	8
PHA-533290 	64	PHA-537098 	16
PHA-537085 	16	PHA-537100 	16

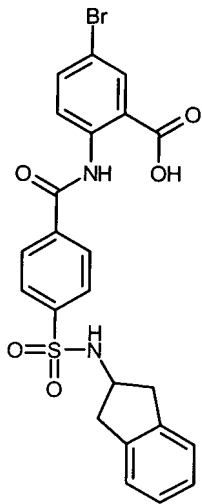
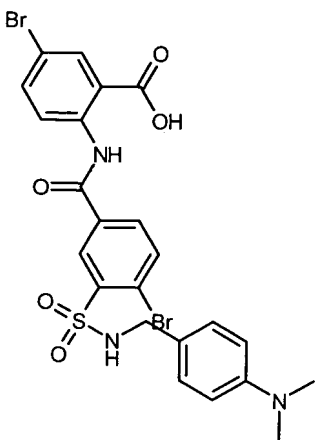
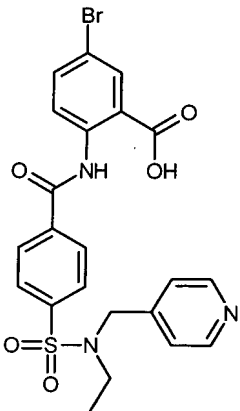
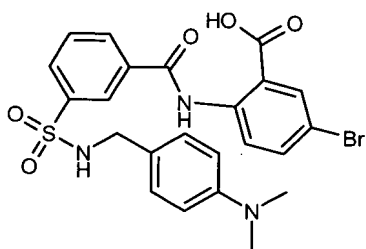
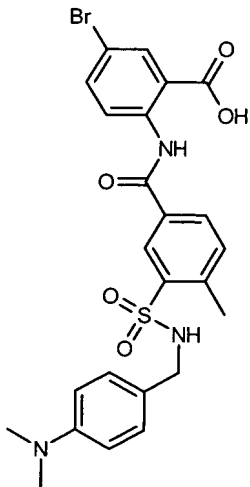
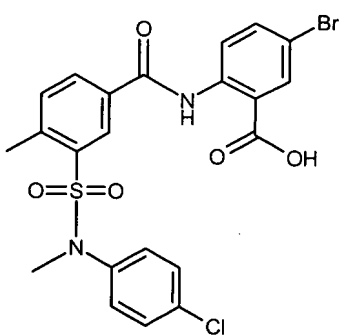
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-537090  least retained isomer by RP-LC/MS	32	PHA-537106 	8
PHA-537092 	16	PHA-537112 	128
PHA-537099 	8	PHA-537121 	4

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-537101 	4	PHA-537128 	8
PHA-537110 	64	PHA-537138 	32
PHA-537114 	16	PHA-537142 	4
PHA-537122 	16	PHA-537144 	8

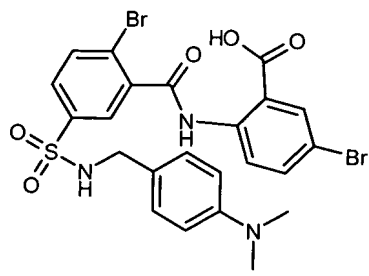
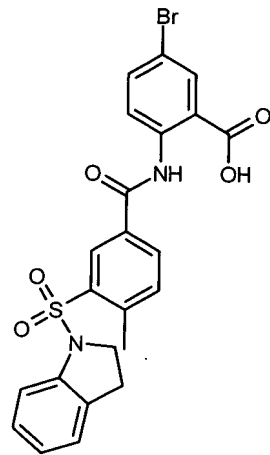
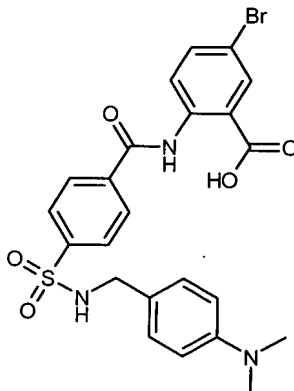
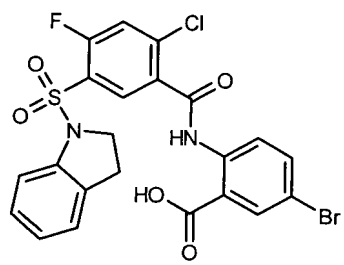
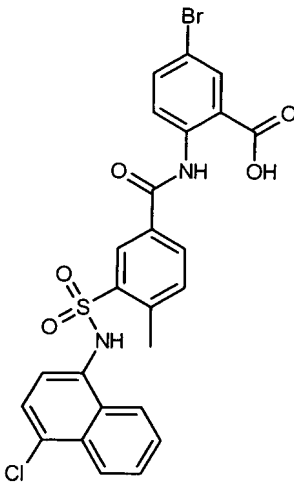
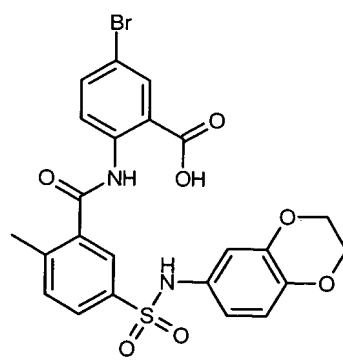
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-537133 	8	PHA-537152 	8
PHA-537139 	4	PHA-537157 	32
PHA-537143 	16	PHA-537162 	16
PHA-537150 	32	PHA-537203  most highly retained isomer by RP-LC/MS	32

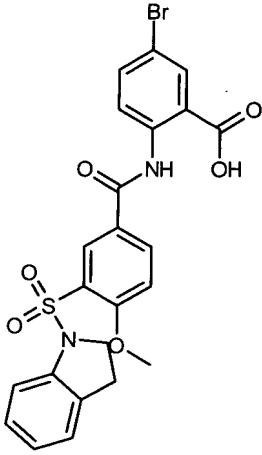
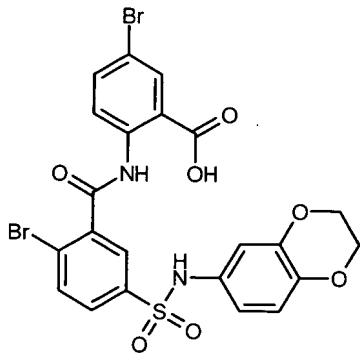
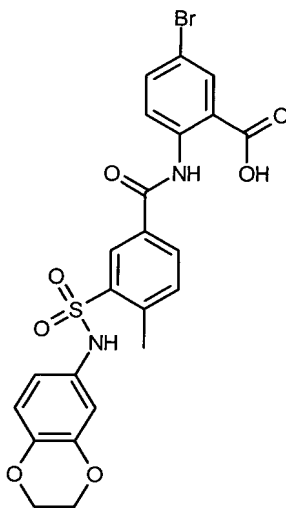
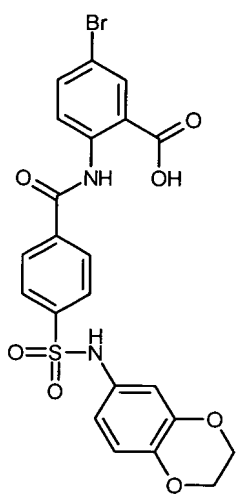
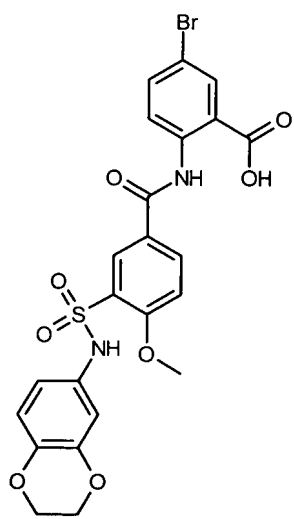
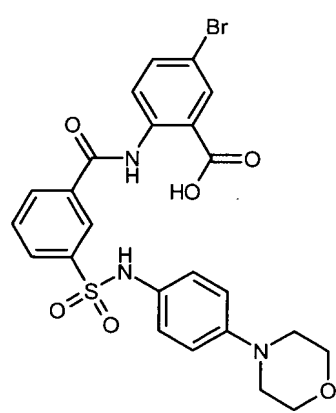
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-537155 	64	PHA-538016 	64
PHA-537158 	32	PHA-539146 	128
PHA-537202  most highly retained isomer by RP-LC/MS	8	PHA-539149 	64

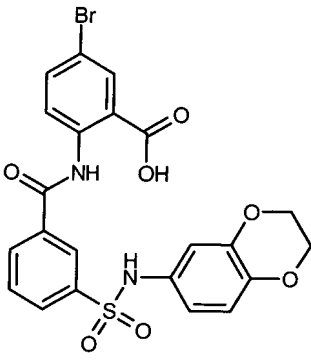
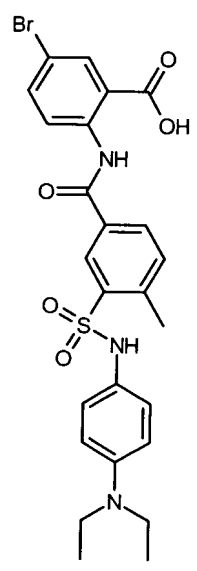
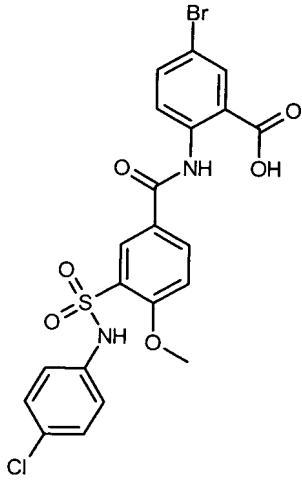
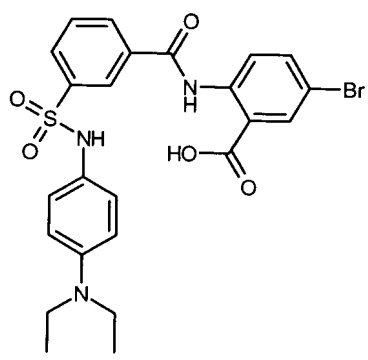
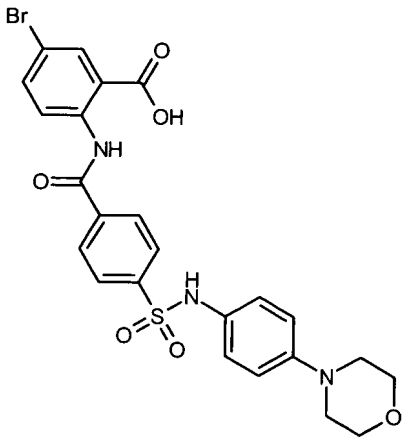
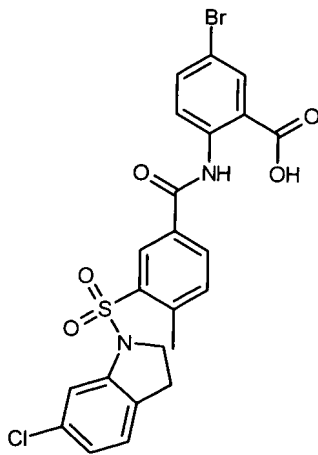
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-537204  most highly retained isomer by RP-LC/MS	64	PHA-539152 	64
PHA-539148 	64	PHA-539154 	32
PHA-539150 	64	PHA-539156 	8

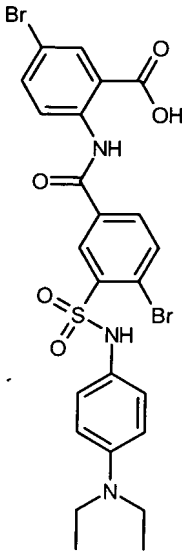
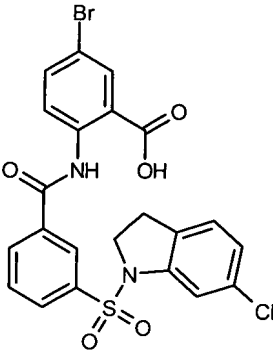
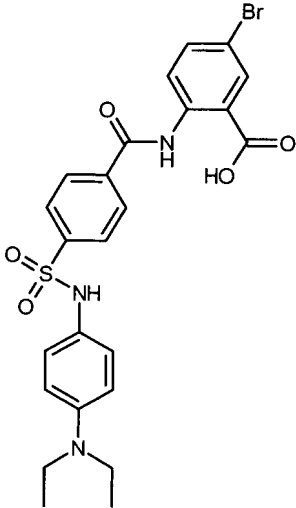
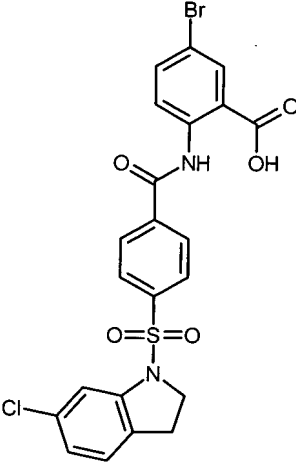
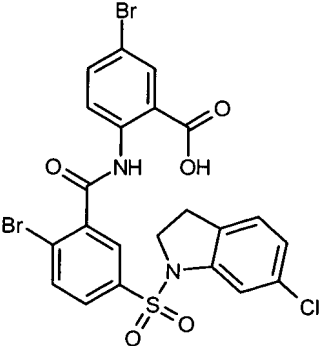
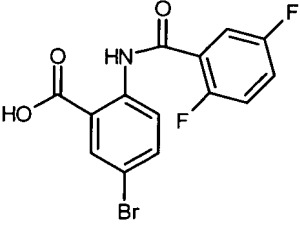
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539153 	32	PHA-539168 	64
PHA-539155 	32	PHA-539170 	64
PHA-539164 	128	PHA-539172 	1

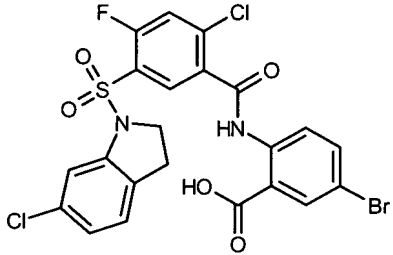
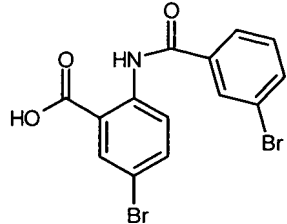
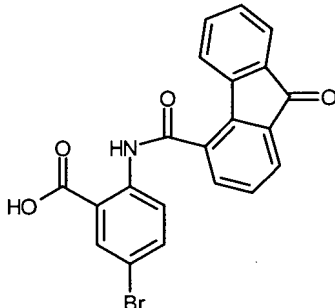
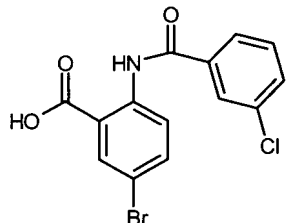
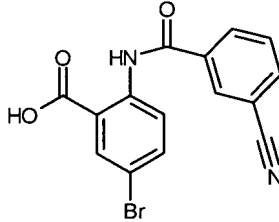
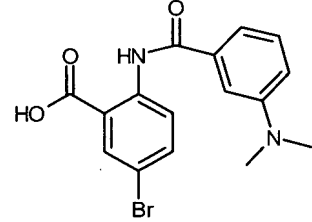
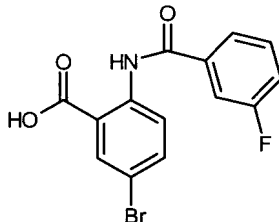
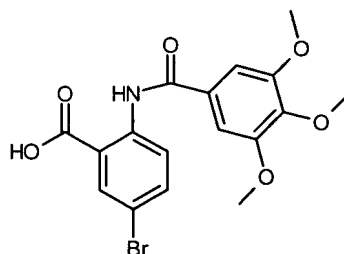
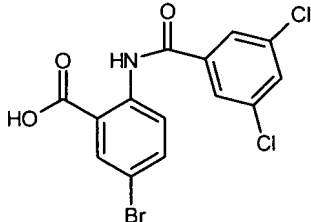
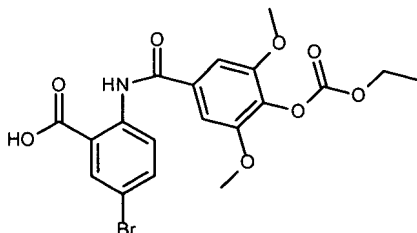


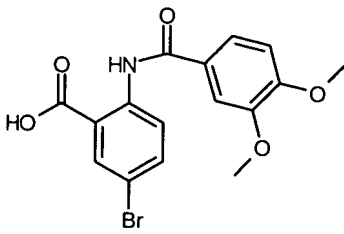
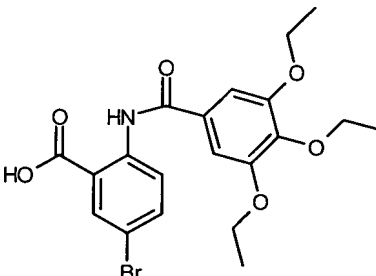
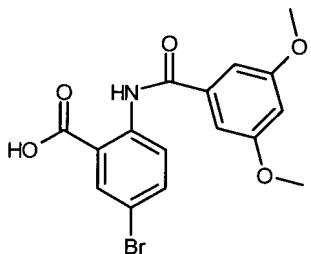
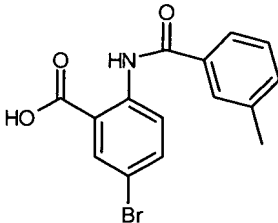
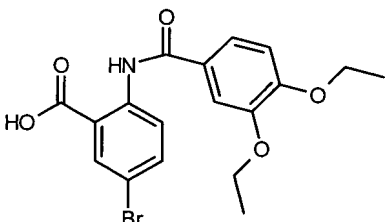
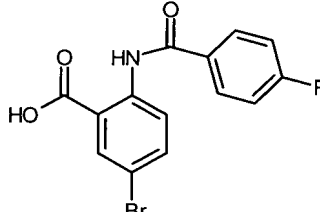
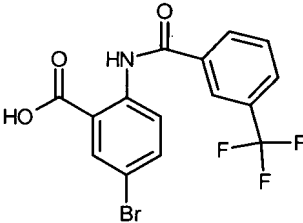
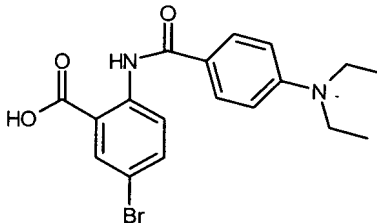
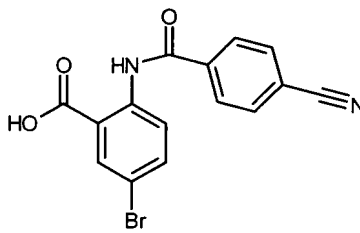
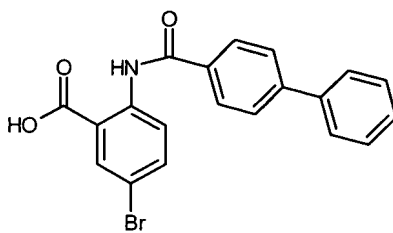
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539169 	16	PHA-539175 	1
PHA-539171 	128	PHA-539179 	8
PHA-539174 	8	PHA-539181 	64

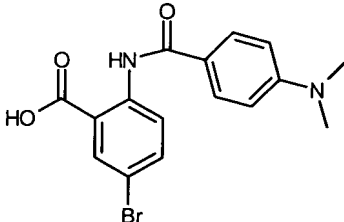
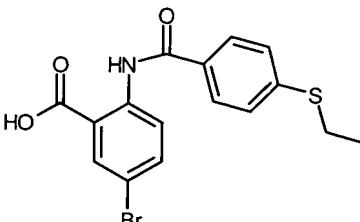
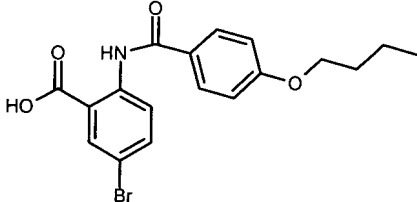
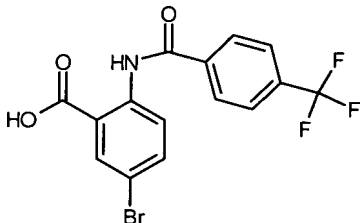
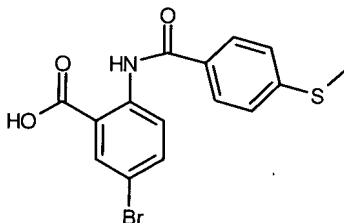
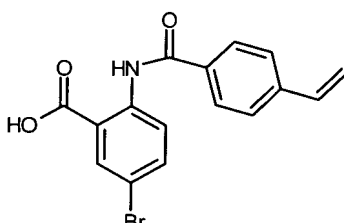
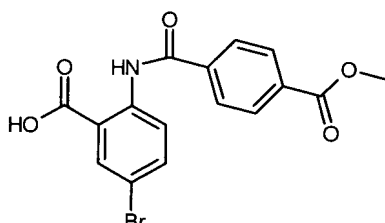
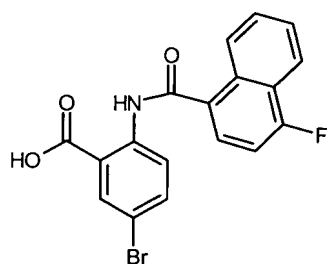
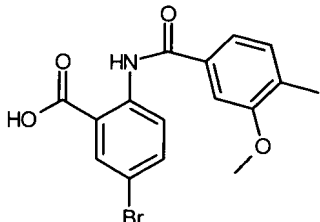
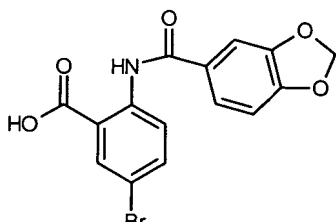
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539177 	4	PHA-539186 	0.5
PHA-539180 	1	PHA-539188 	16
PHA-539183 	8	PHA-539193 	32

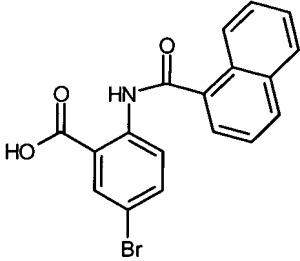
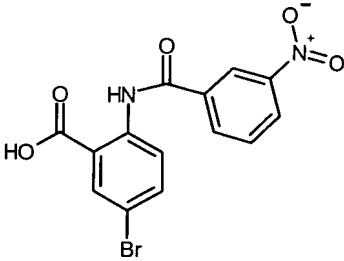
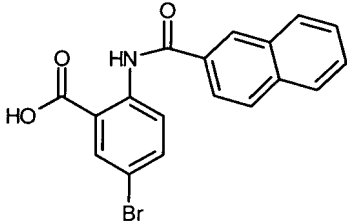
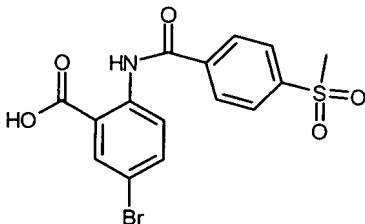
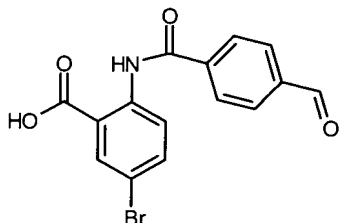
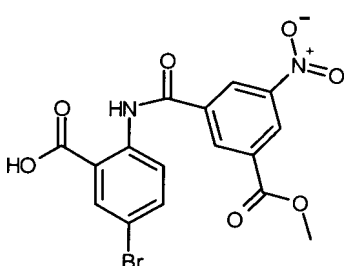
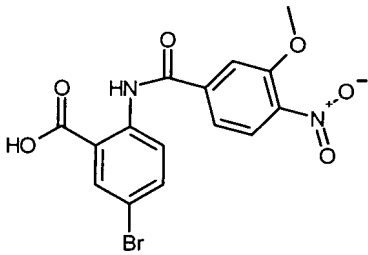
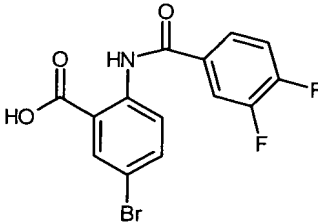
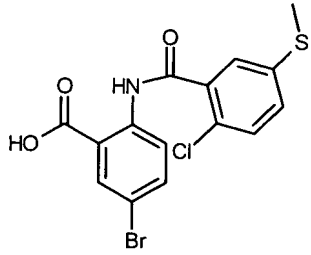
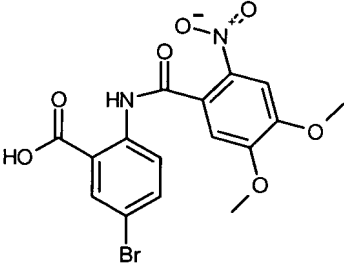
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539187 	4	PHA-539195 	64
PHA-539190 	16	PHA-539198 	128
PHA-539194 	64	PHA-539203 	1

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539197 	64	PHA-539207 	2
PHA-539199 	32	PHA-539209 	0.5
PHA-539206 	2	PHA-539235 	128

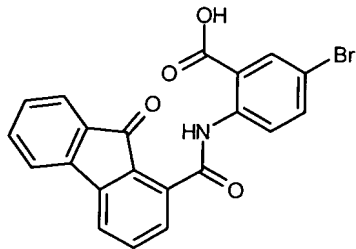
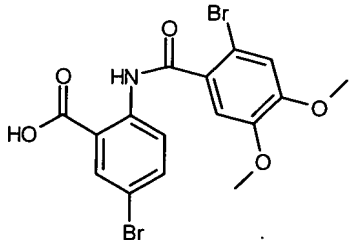
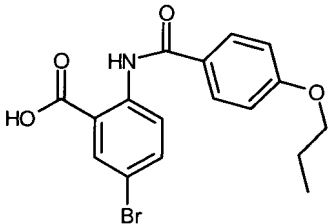
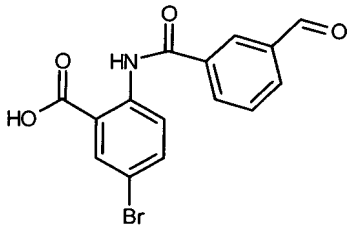
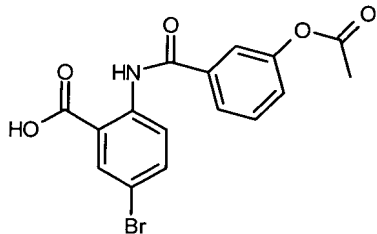
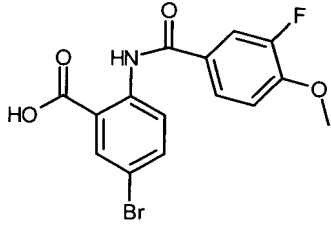
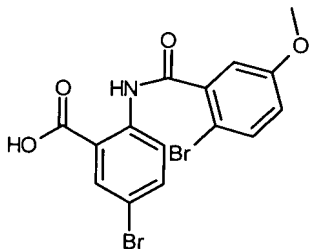
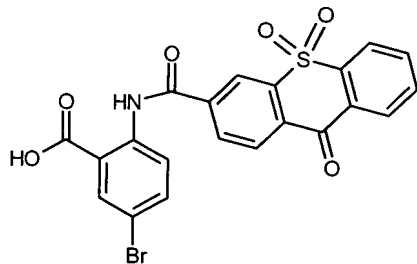
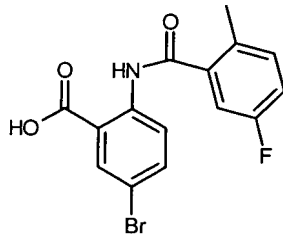
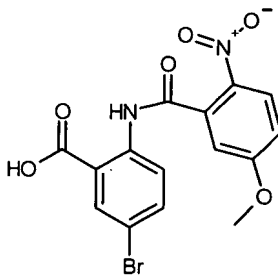
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539208 	16	PHA-539246 	8
PHA-539234 	128	PHA-539248 	8
PHA-539245 	>128	PHA-539250 	32
PHA-539247 	64	PHA-539252 	32
PHA-539249 	8	PHA-539254 	128

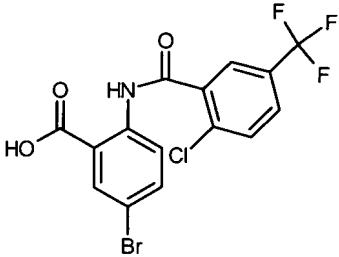
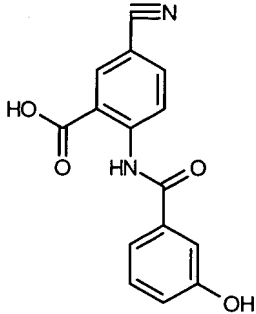
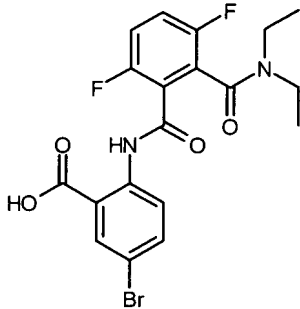
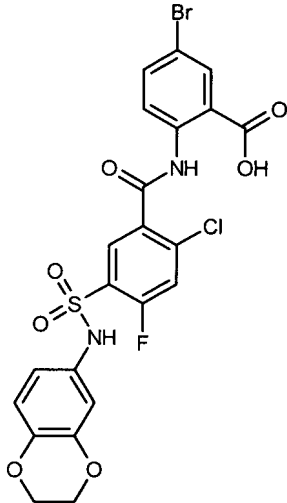
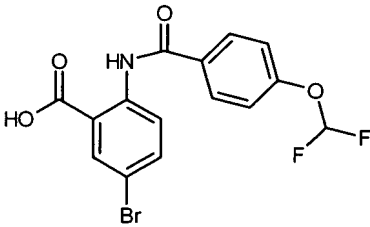
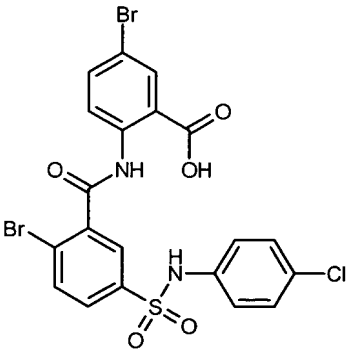
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539251 	128	PHA-539256 	32
PHA-539253 	16	PHA-539258 	64
PHA-539255 	>128	PHA-539260 	64
PHA-539257 	8	PHA-539263 	32
PHA-539259 	128	PHA-539265 	32

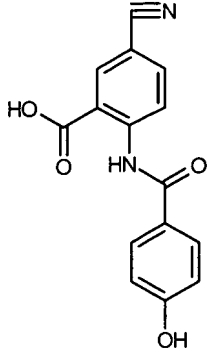
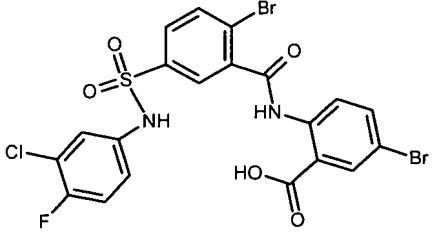
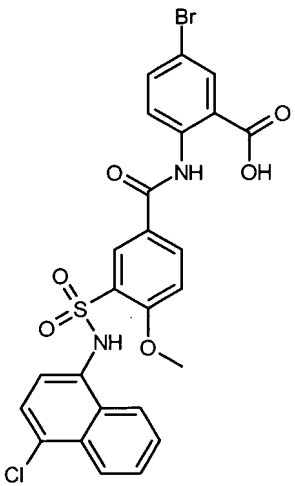
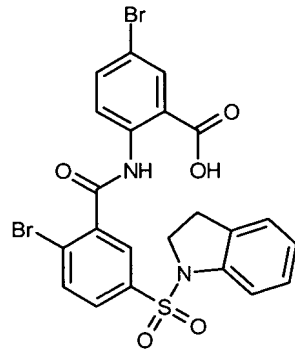
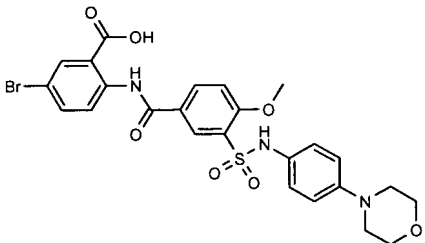
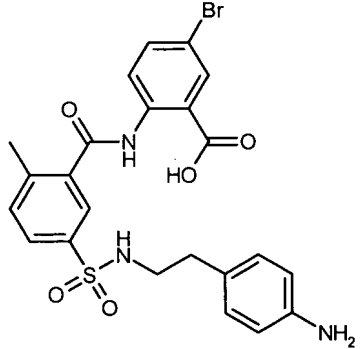
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539262 	32	PHA-539267 	0.5
PHA-539264 	8	PHA-539269 	32
PHA-539266 	2	PHA-539271 	>128
PHA-539268 	32	PHA-539277 	32
PHA-539270 	>128	PHA-539285 	16

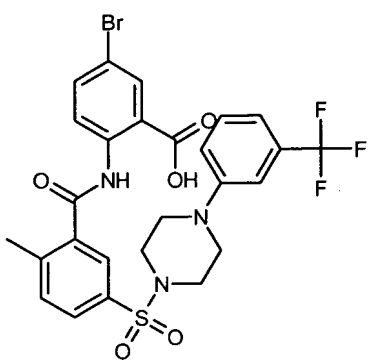
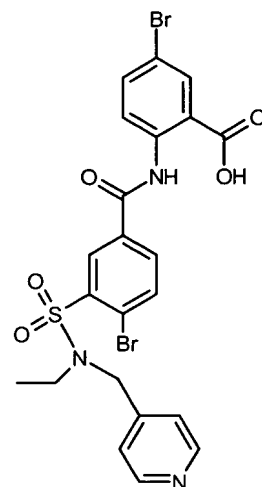
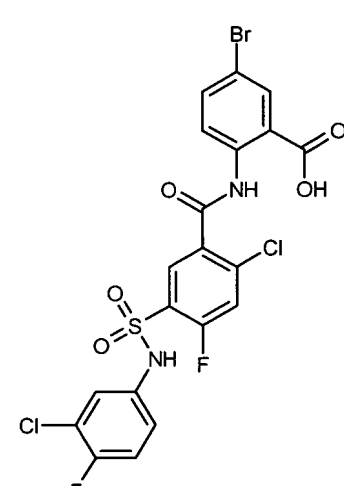
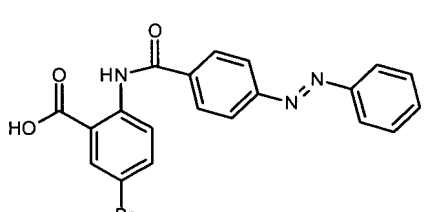
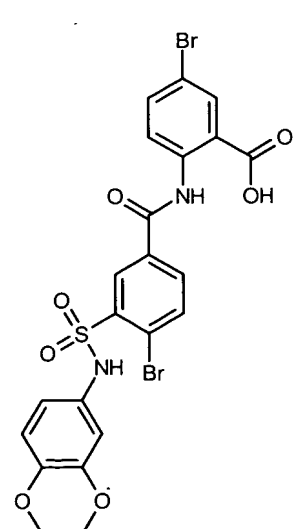
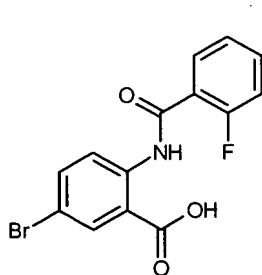
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539276 	32	PHA-539294 	128
PHA-539278 	2	PHA-539296 	64
PHA-539293 	>128	PHA-539298 	64
PHA-539295 	32	PHA-539303 	32
PHA-539297 	>128	PHA-539307 	>128

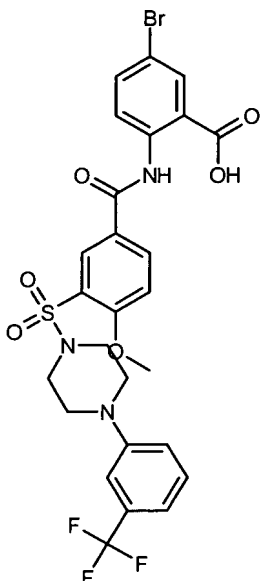
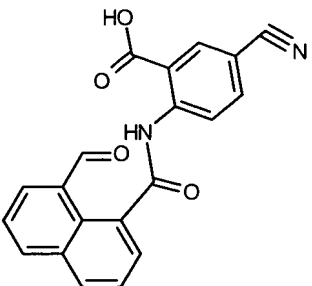
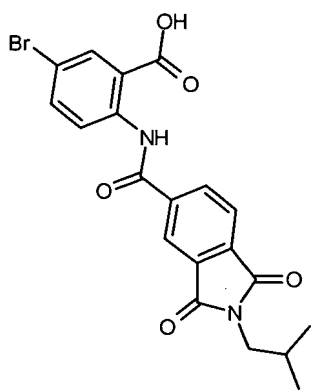
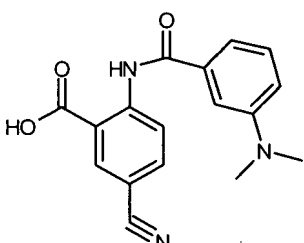
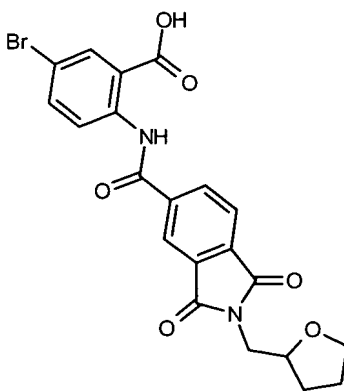
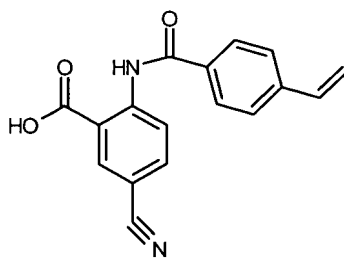


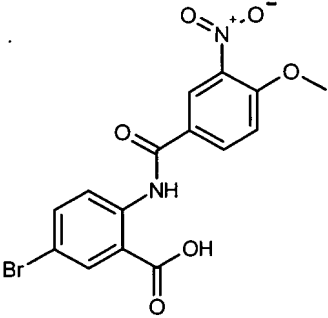
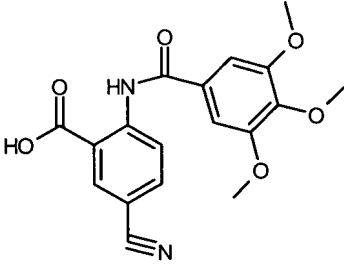
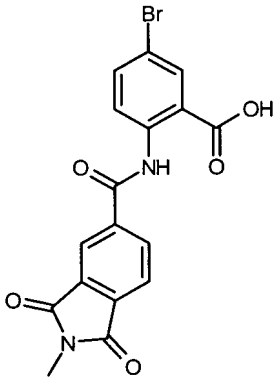
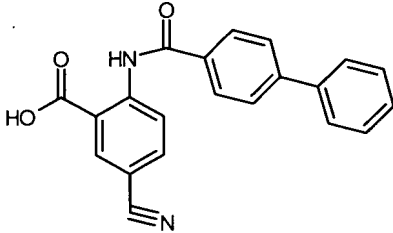
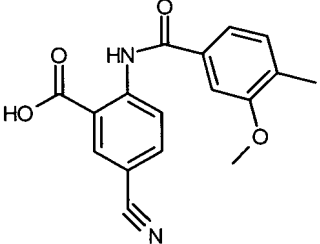
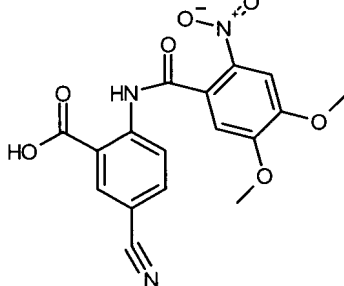
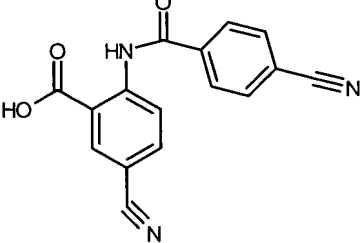
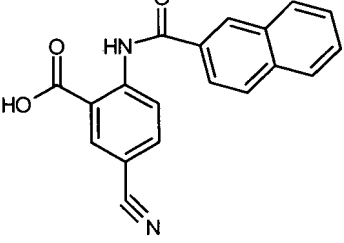
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539302 	>128	PHA-539310 	>128
PHA-539305 	64	PHA-539313 	>128
PHA-539308 	128	PHA-539317 	16
PHA-539312 	128	PHA-539322 	16
PHA-539314 	64	PHA-539329 	>128

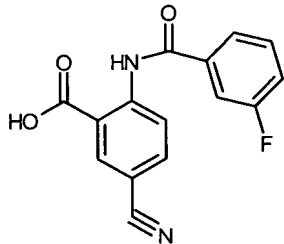
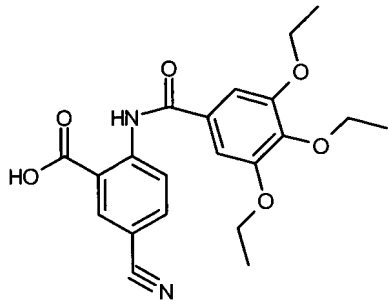
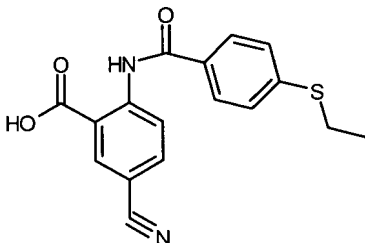
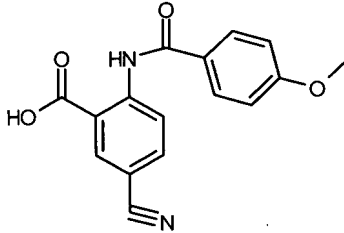
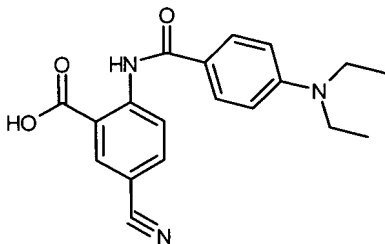
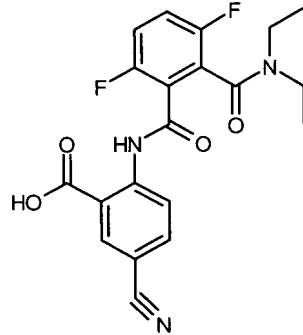
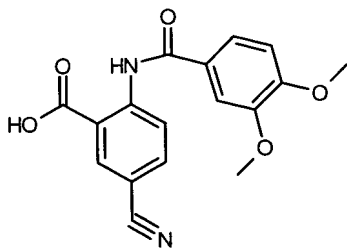
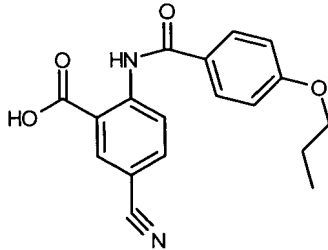
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539318 	>128	PHA-539337 	32
PHA-539328 	>128	PHA-543684 	128
PHA-539332 	64	PHA-543686 	4

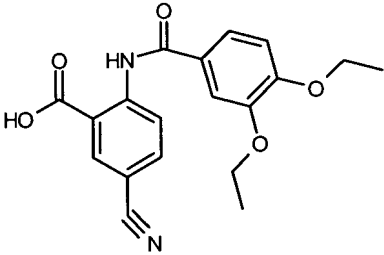
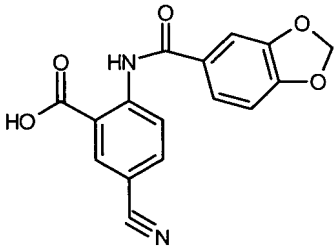
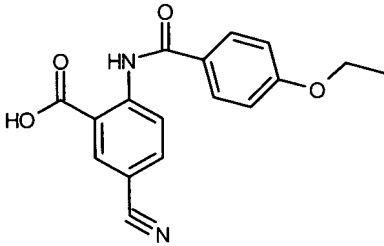
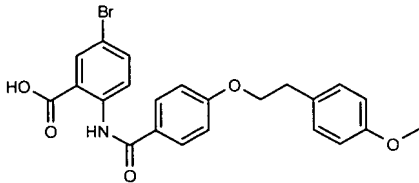
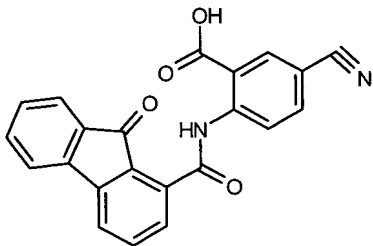
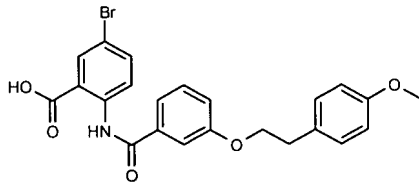
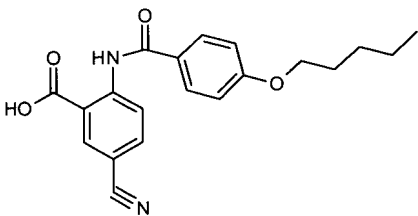
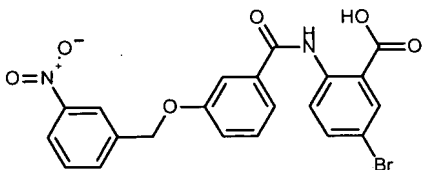
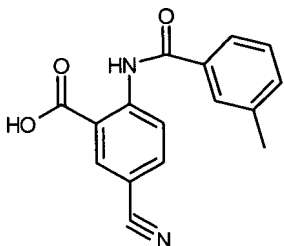
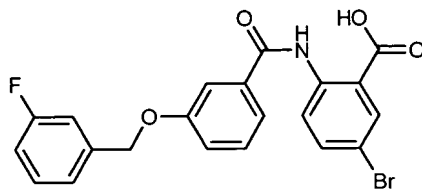
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-539338 	64	PHA-543690 	32
PHA-543685 	>128	PHA-543693 	2
PHA-543689 	64	PHA-543698 	>128

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-543692 	16	PHA-543701 	128
PHA-543695 	>128	PHA-543708 	16
PHA-543700 	64	PHA-551716 	128

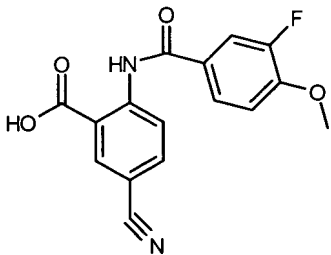
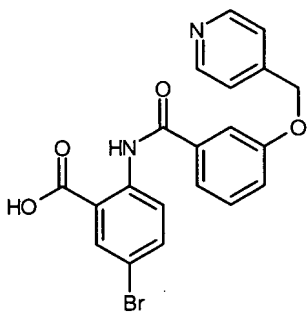
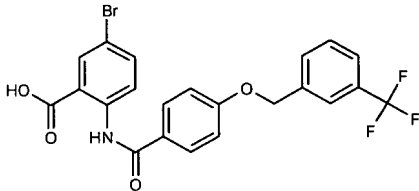
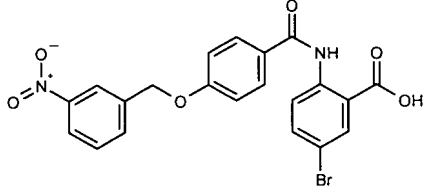
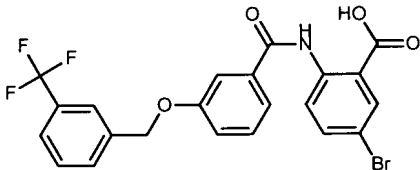
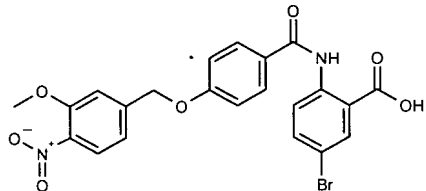
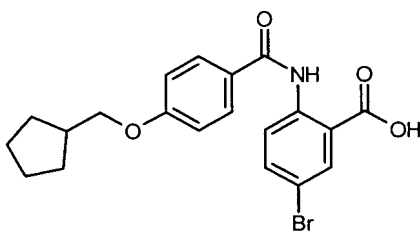
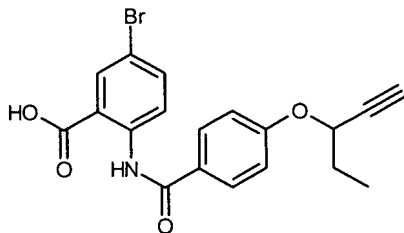
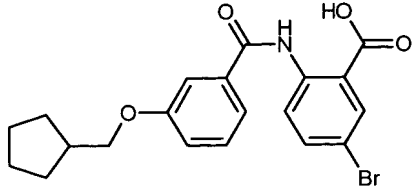
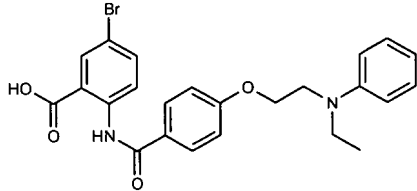
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-543706 	32	PHA-563331 	>128
PHA-551625 	2	PHA-563335 	8
PHA-551672 	8	PHA-563341 	8

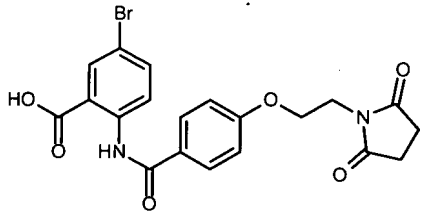
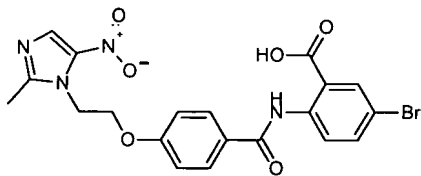
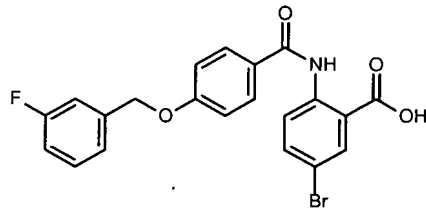
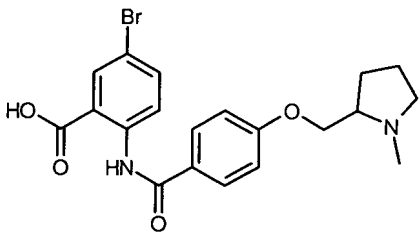
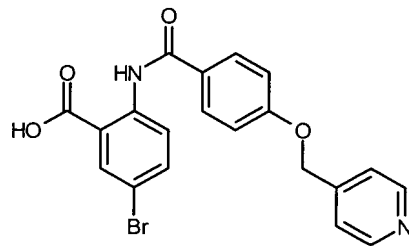
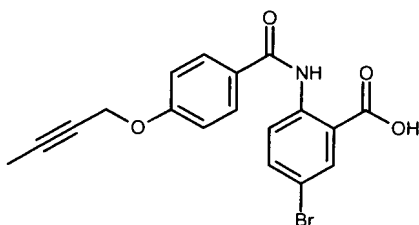
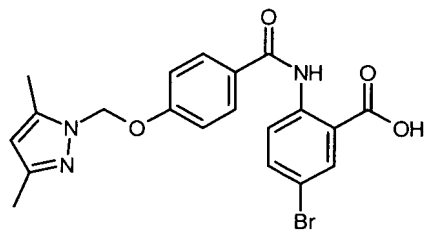
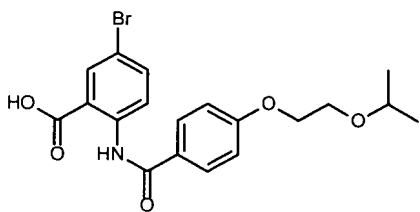
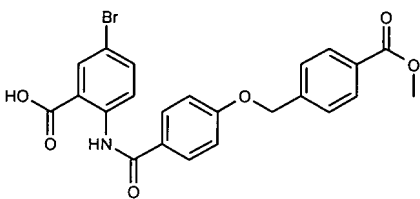
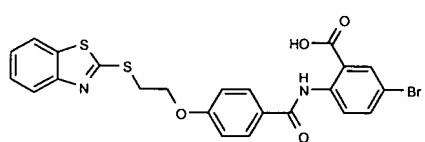
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-551675 	32	PHA-563344 	64
PHA-556420 	128	PHA-563347 	64
PHA-563330 	>128	PHA-563351 	>128
PHA-563333 	>128	PHA-563354 	2

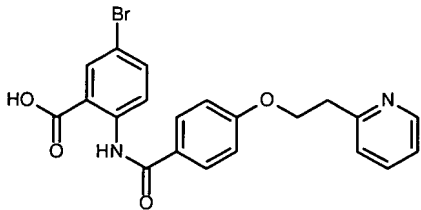
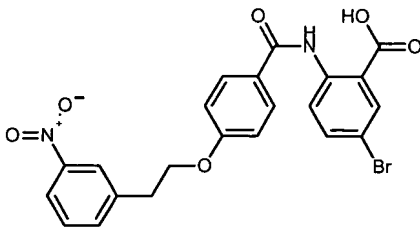
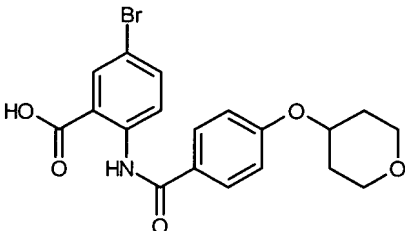
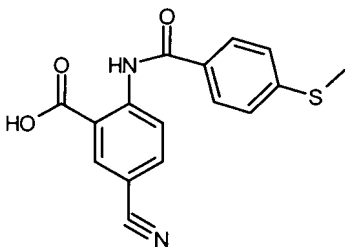
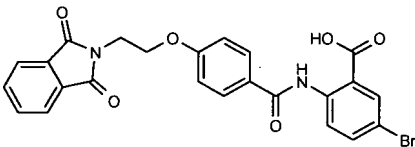
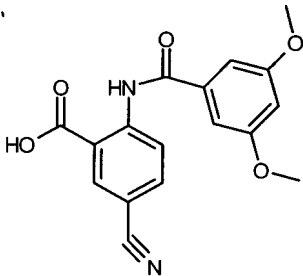
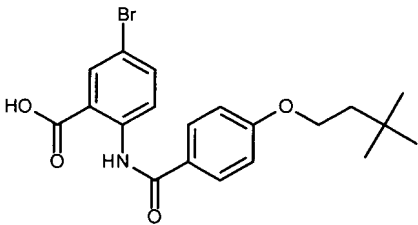
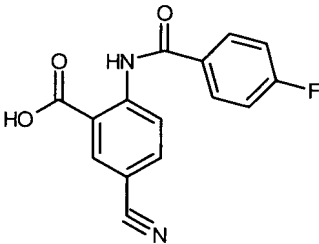
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563340 	64	PHA-563363 	16
PHA-563342 	2	PHA-563365 	16
PHA-563345 	16	PHA-563368 	>128
PHA-563350 	64	PHA-563371 	16

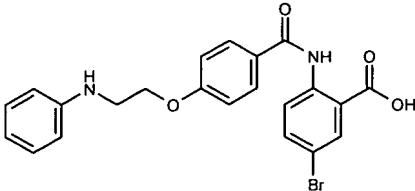
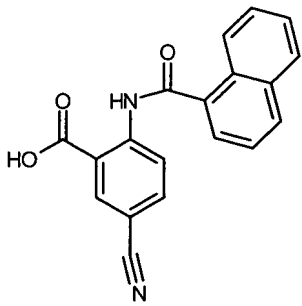
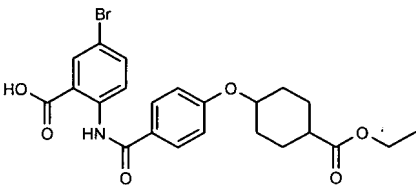
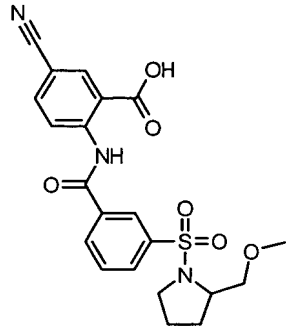
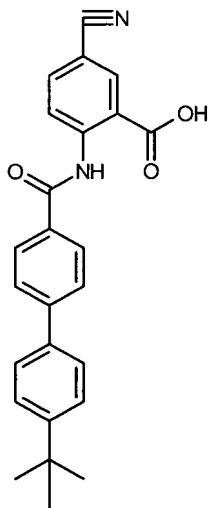
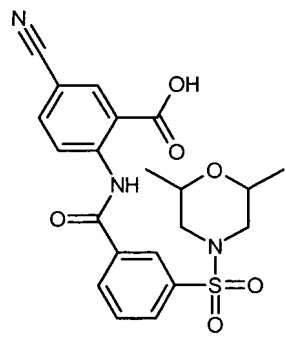
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563353 	128	PHA-563378 	16
PHA-563360 	32	PHA-563388 	>128
PHA-563364 	>128	PHA-563390 	32
PHA-563366 	4	PHA-563392 	16
PHA-563370 	32	PHA-563394 	16

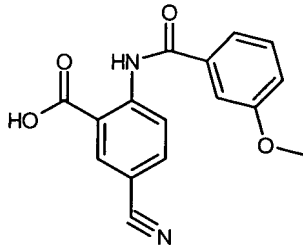
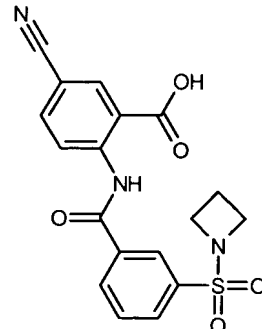
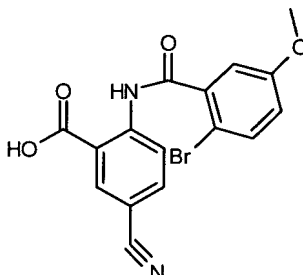
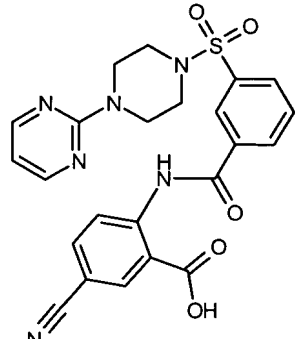
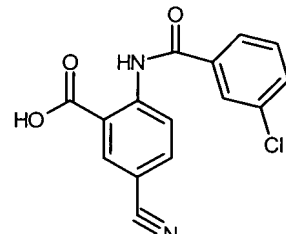
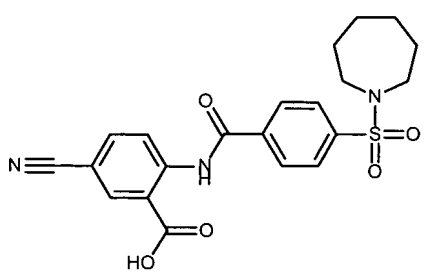
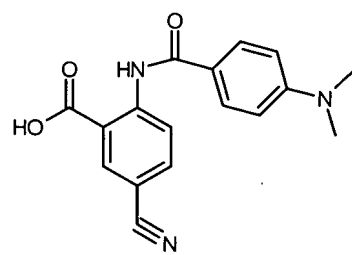
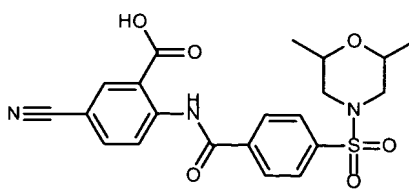


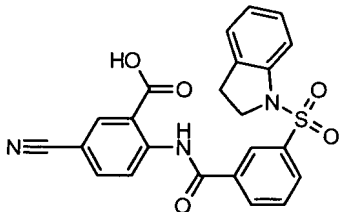
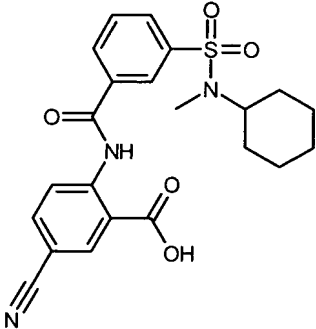
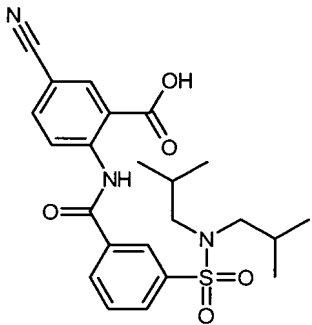
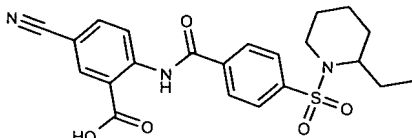
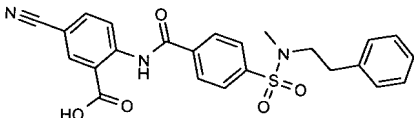
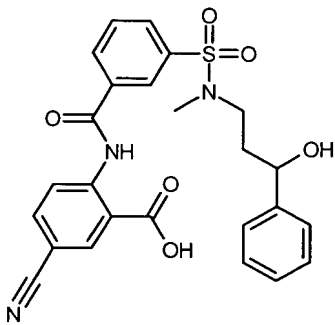
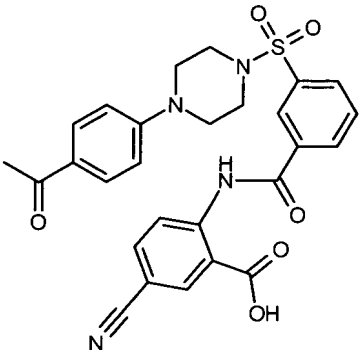
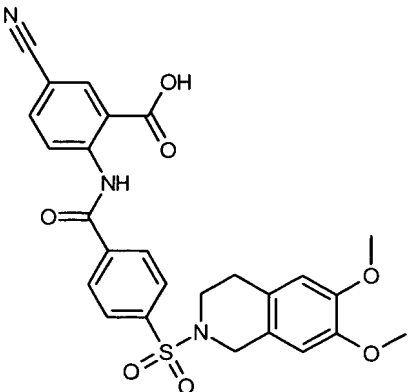
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563375 	8	PHA-563398 	>128
PHA-563386 	32	PHA-563399 	16
PHA-563389 	64	PHA-563404 	8
PHA-563391 	>128	PHA-563407 	>128
PHA-563393 	128	PHA-563409 	64

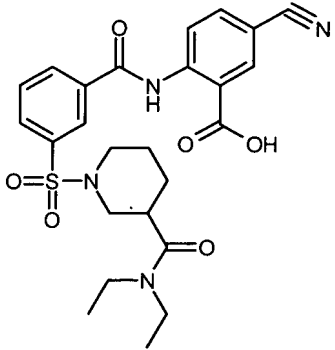
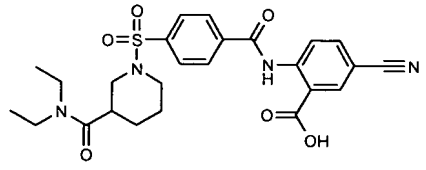
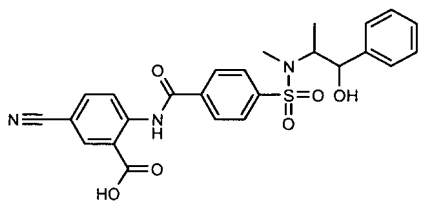
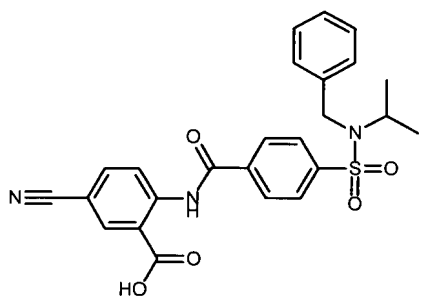
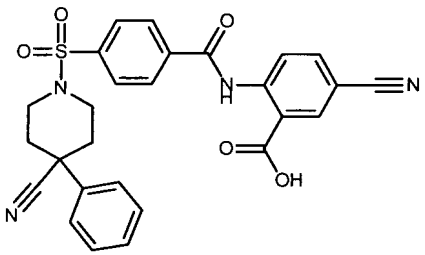
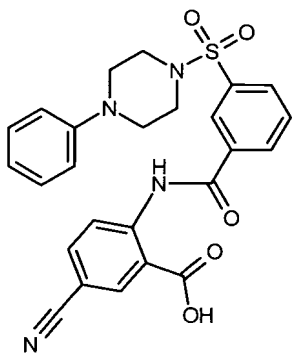
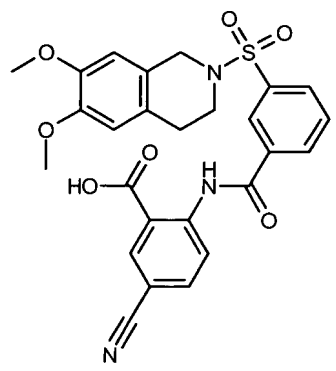
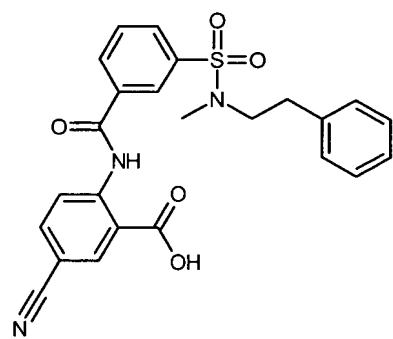
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563396 	>128	PHA-563413 	128
PHA-563397 	32	PHA-563417 	>128
PHA-563401 	>128	PHA-563420 	16
PHA-563406 	64	PHA-563427 	>128
PHA-563408 	>128	PHA-563441 	64

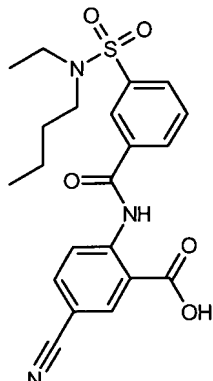
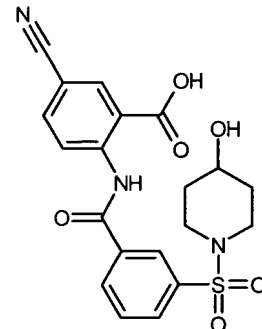
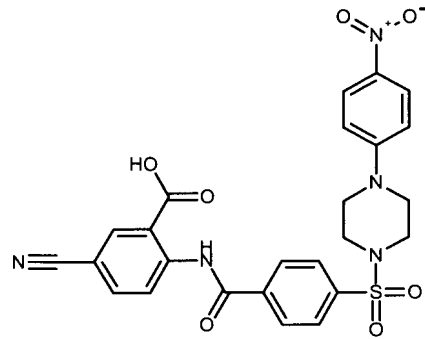
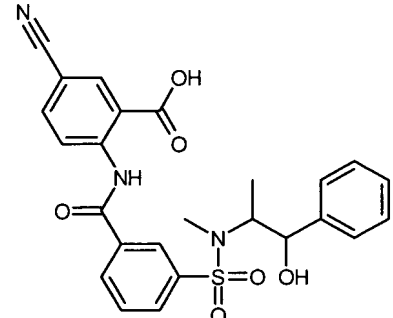
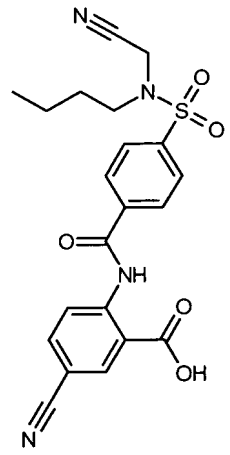
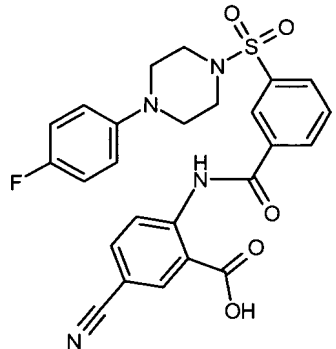
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563411 	128	PHA-563449 	64
PHA-563415 	128	PHA-571150 	0.5
PHA-563419 	64	PHA-571152 	8
PHA-563426 	64	PHA-571154 	128

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-563440 	64	PHA-571156 	16
PHA-563442 	>128	PHA-571160 	64
PHA-569976 	32	PHA-571162 	16

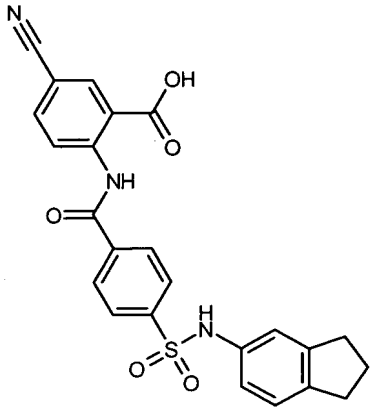
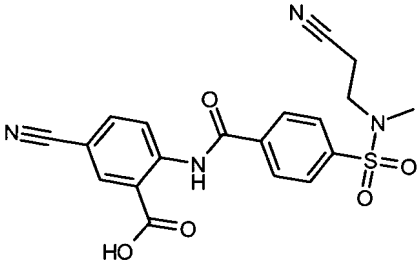
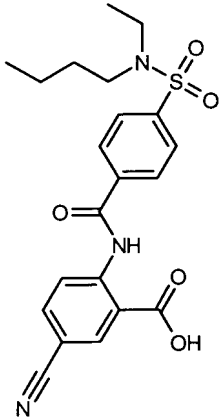
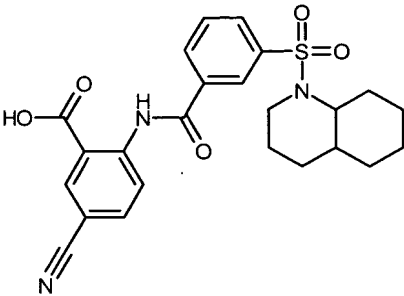
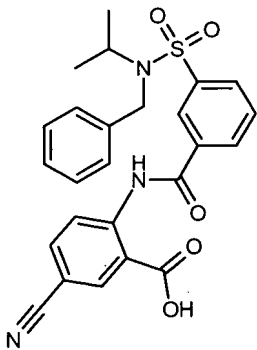
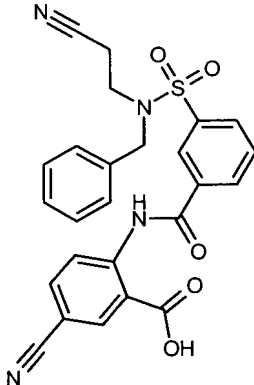
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571151 	8	PHA-571167 	32
PHA-571153 	64	PHA-571170 	64
PHA-571155 	32	PHA-571174 	64
PHA-571157 	32	PHA-571182 	64

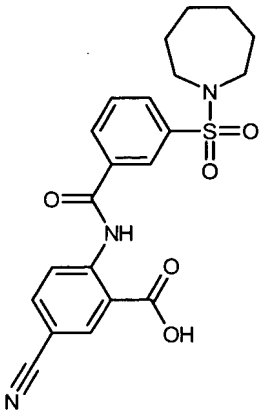
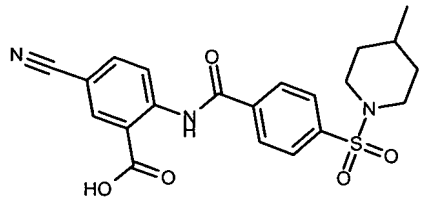
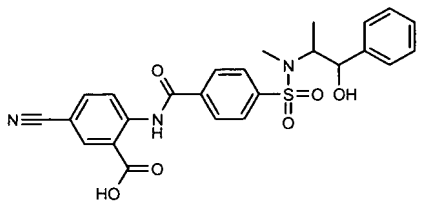
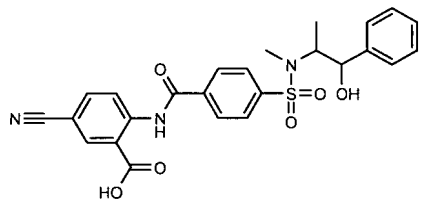
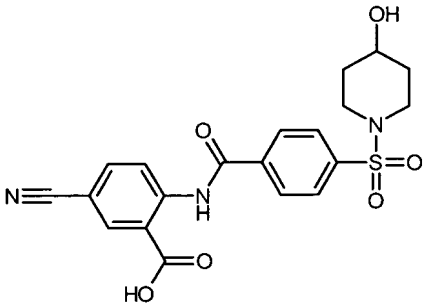
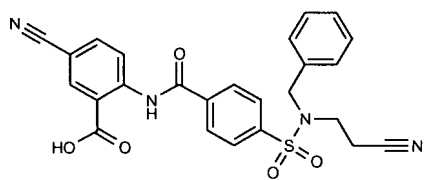
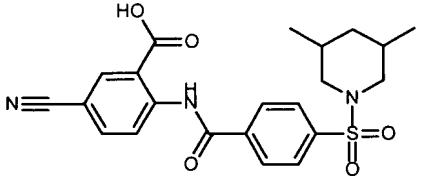
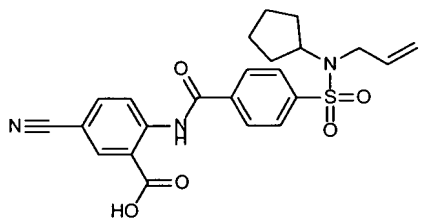
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571161 	>128	PHA-571186 	128
PHA-571164 	8	PHA-571189 	64
PHA-571169 	32	PHA-571196 	64
PHA-571172 	32	PHA-571198 	>128

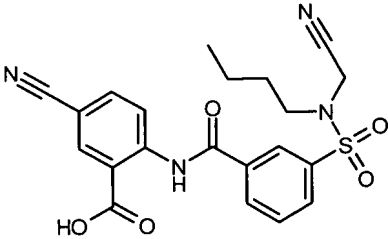
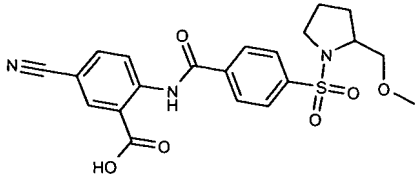
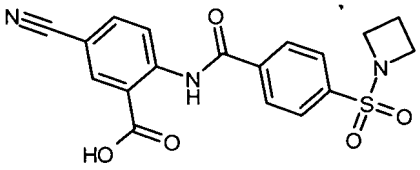
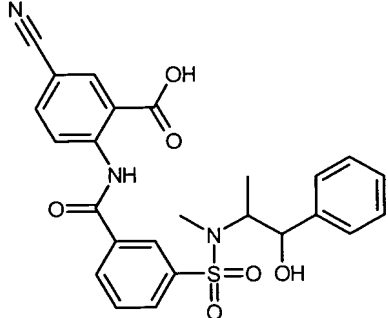
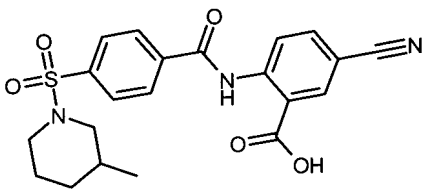
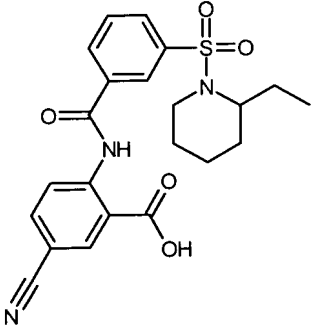
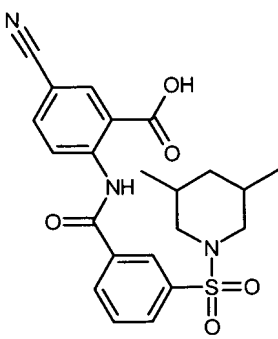
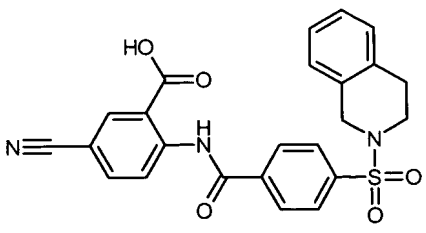
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571176 	64	PHA-571202 	128
PHA-571183 	32	PHA-571205 	32
PHA-571188 	8	PHA-571208 	64
PHA-571194 	4	PHA-571215 	8

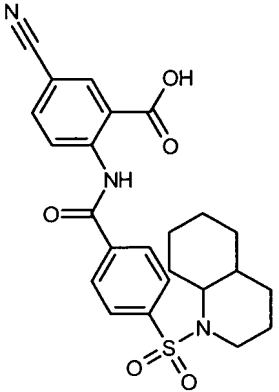
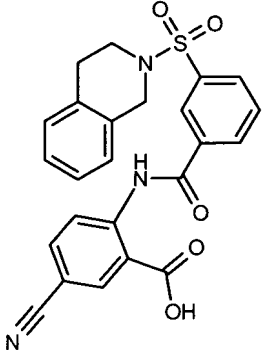
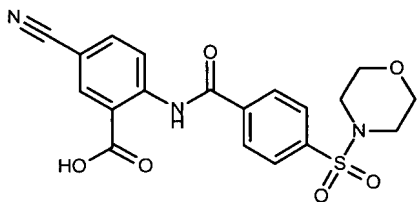
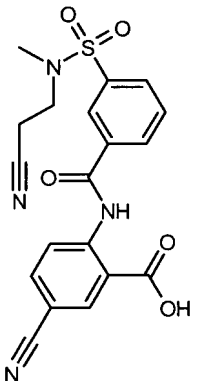
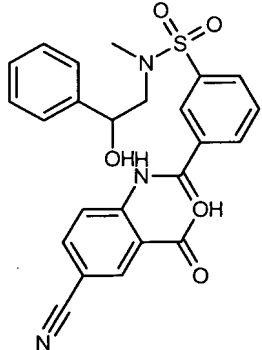
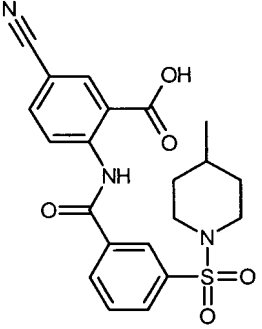
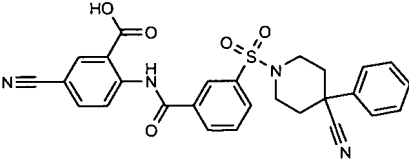
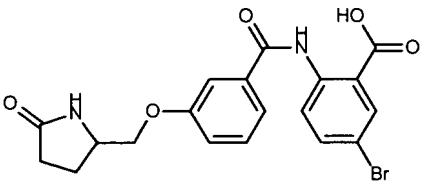
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571197 	16	PHA-571219 	32
PHA-571199 	64	PHA-571226 	64
PHA-571203 	32	PHA-571230 	16

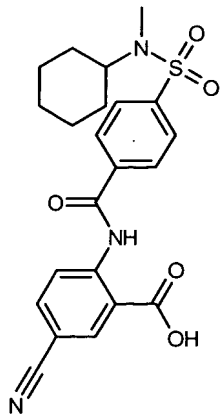
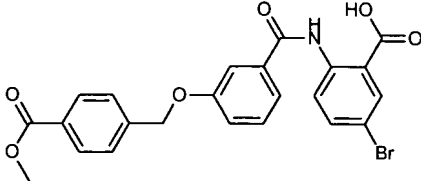
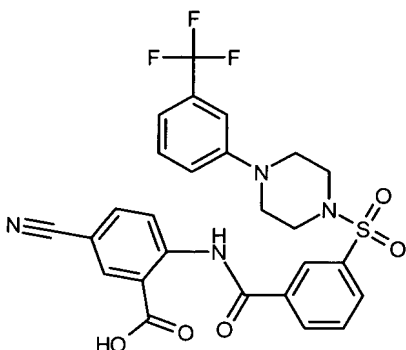
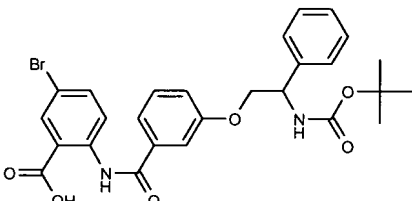
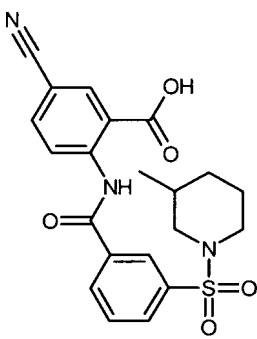
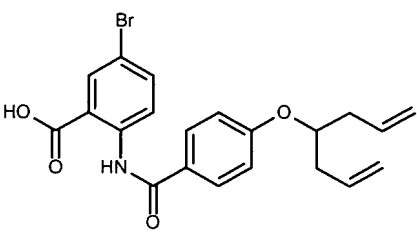


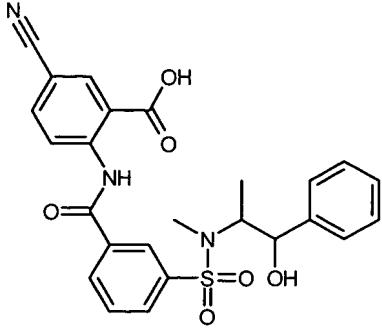
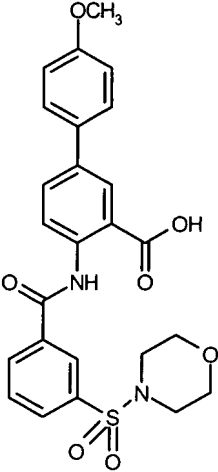
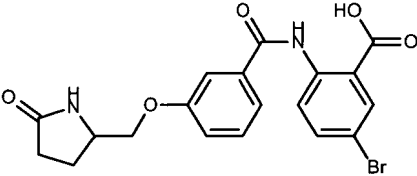
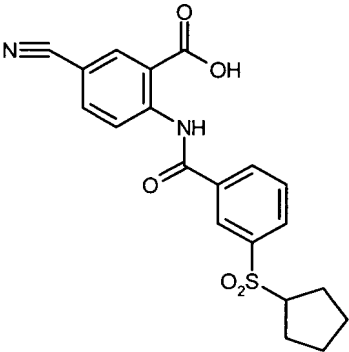
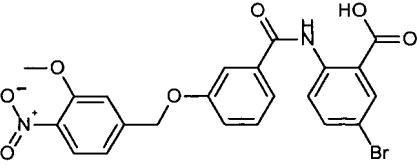
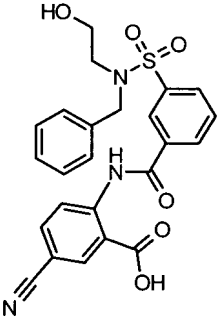
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571207 	32	PHA-571232 	>128
PHA-571214 	16	PHA-571235 	8
PHA-571216 	32	PHA-571238 	128

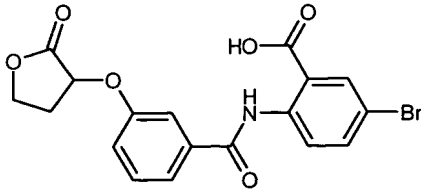
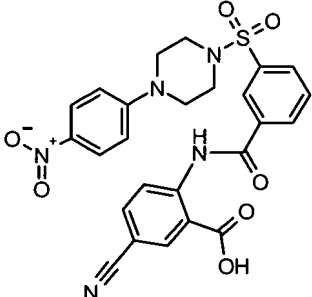
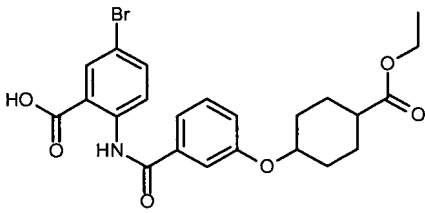
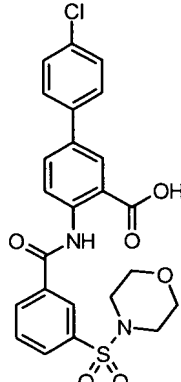
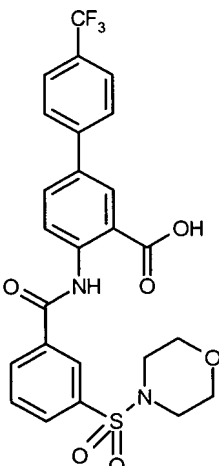
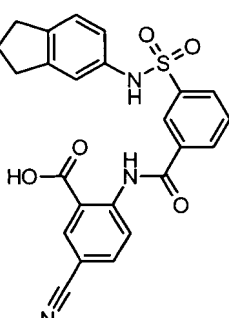
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571224 	8	PHA-571240 	16
PHA-571228 	32	PHA-571242 	32
PHA-571231 	>128	PHA-571246 	32
PHA-571234 	8	PHA-571253 	16

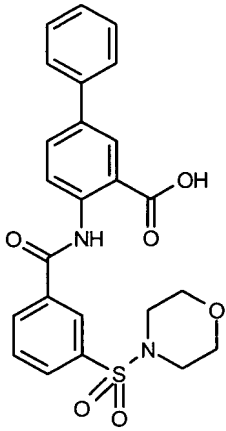
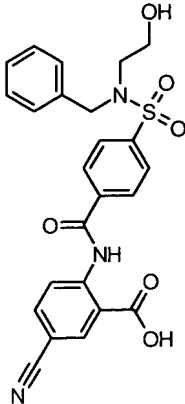
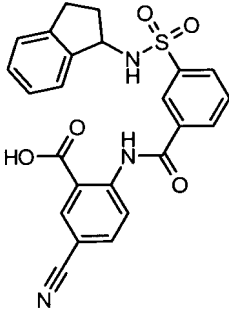
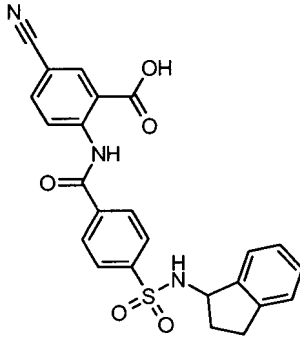
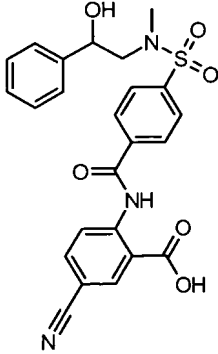
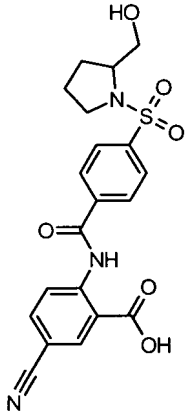
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571237 	16	PHA-571257 	64
PHA-571239 	128	PHA-571260 	32
PHA-571241 	16	PHA-571263 	16
PHA-571243 	4	PHA-571265 	16

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571249 	16	PHA-571269 	16
PHA-571255 	>128	PHA-571271 	64
PHA-571258 	8	PHA-571273 	8
PHA-571262 	32	PHA-571281 	128

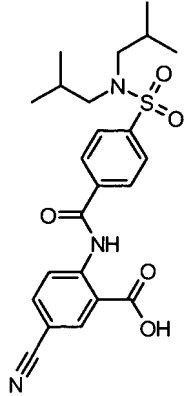
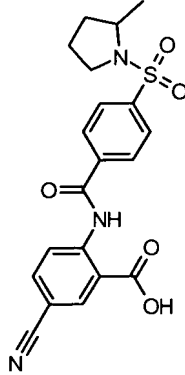
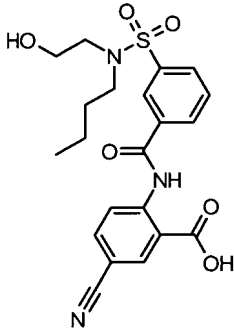
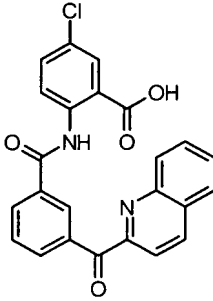
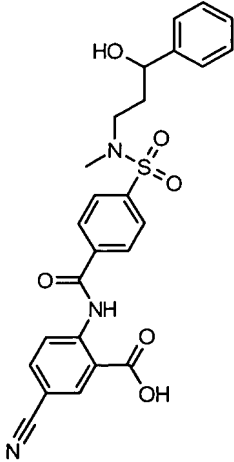
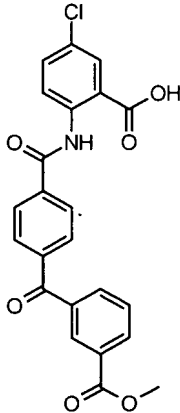
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571264 	32	PHA-571283 	16
PHA-571267 	32	PHA-571287 	2
PHA-571270 	8	PHA-571292 	32

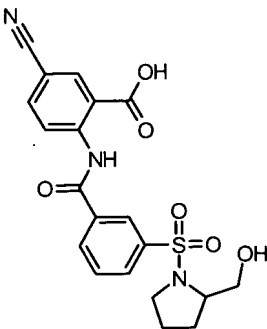
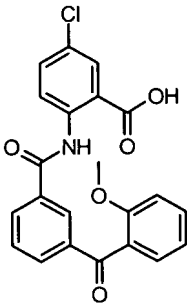
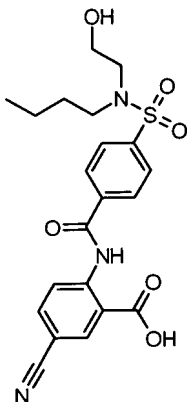
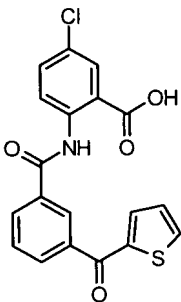
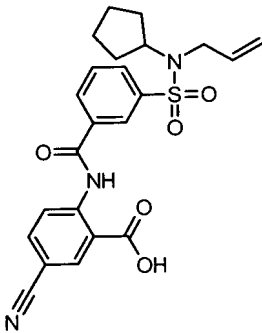
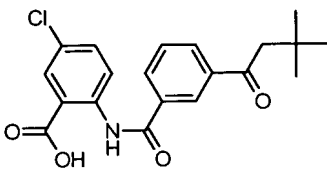
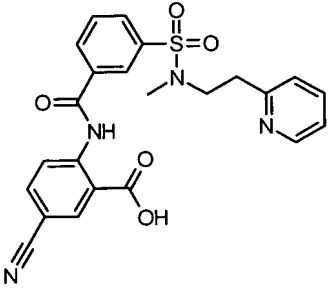
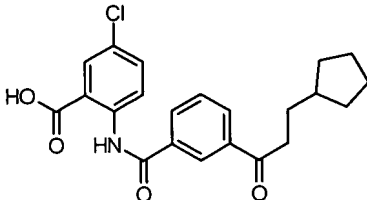
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571272 	32	PHA-610941 	>128
PHA-571280 	>128	PHA-630426 	>128
PHA-571282 	16	PHA-656808 	64

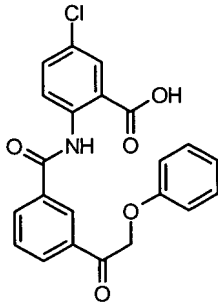
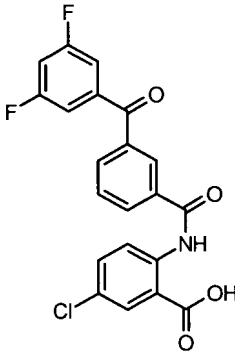
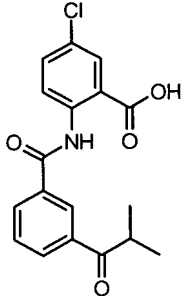
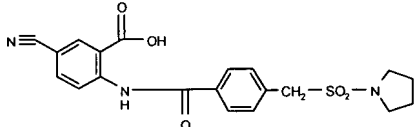
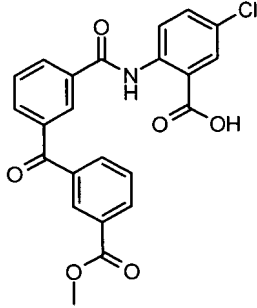
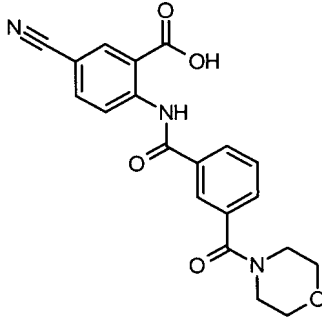
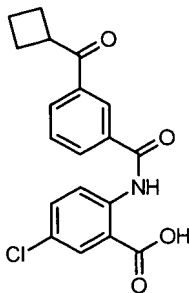
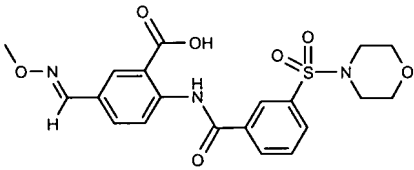
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-571285 	64	PHA-656810 	2
PHA-571289 	32	PHA-656820 	>128
PHA-610940 	>128	PHA-656860 	8

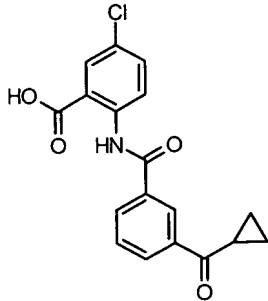
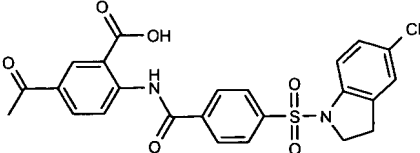
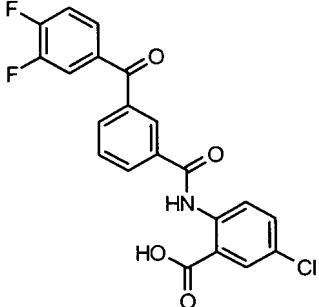
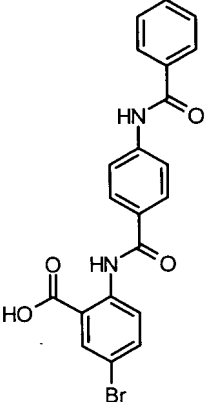
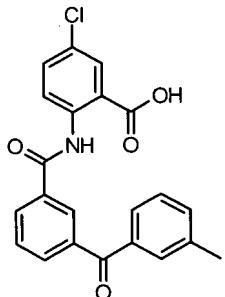
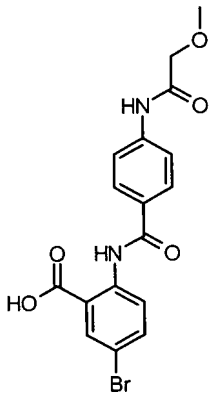
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-610942 	>128	PHA-656862 	32
PHA-656807 	64	PHA-656866 	>128
PHA-656809 	64	PHA-656868 	>128

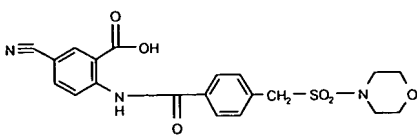
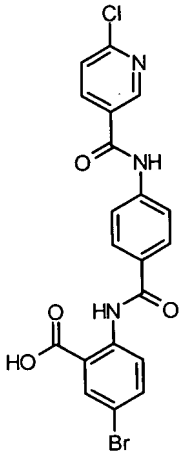
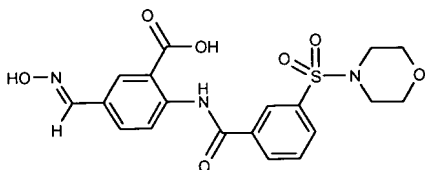
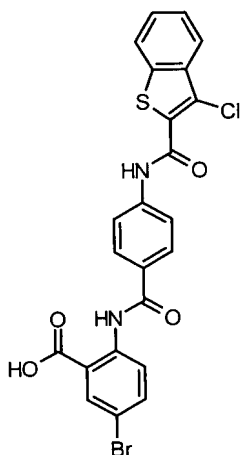
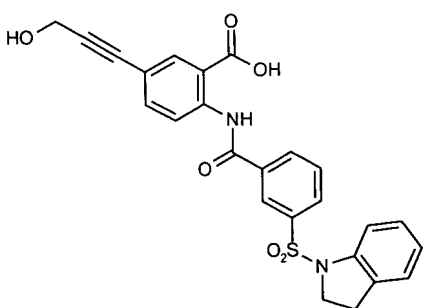
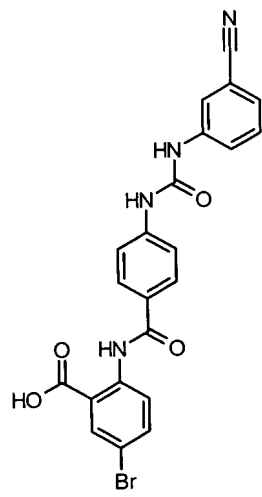


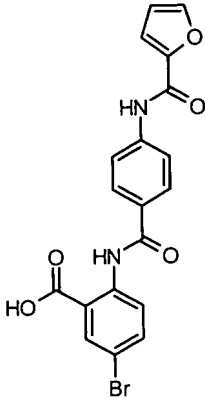
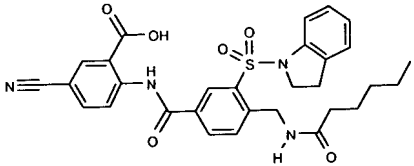
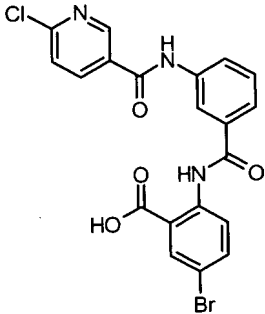
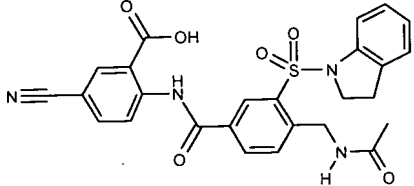
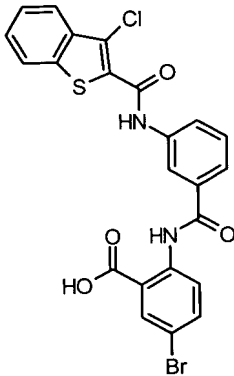
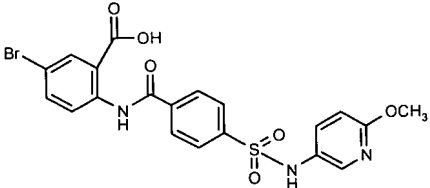
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-656811 	32	PHA-656871 	128
PHA-656859 	16	PHA-656880 	16
PHA-656861 	32	PHA-656883 	16

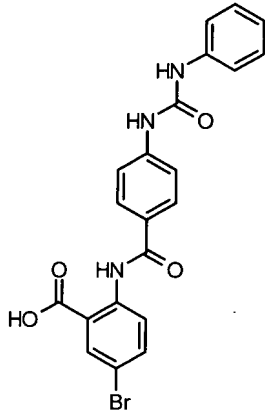
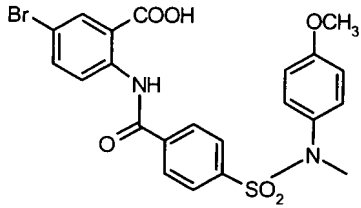
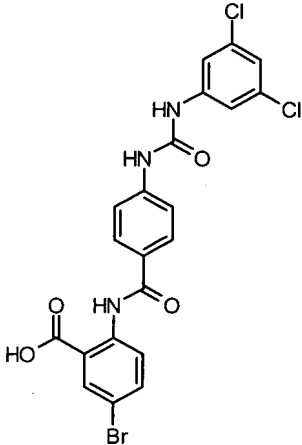
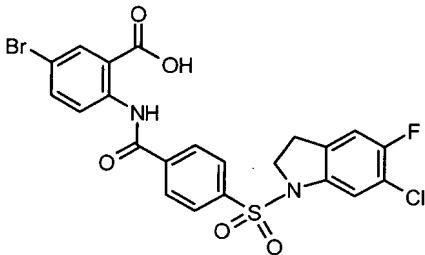
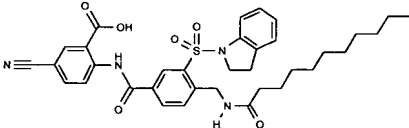
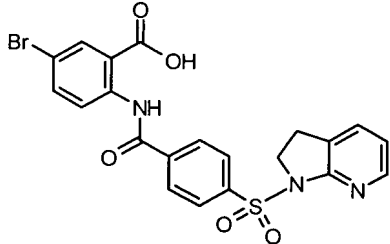
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-656863 	8	PHA-656885 	16
PHA-656867 	64	PHA-656887 	8
PHA-656870 	8	PHA-656889 	16
PHA-656872 	>128	PHA-656891 	16

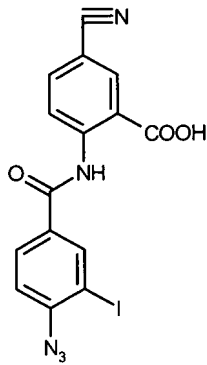
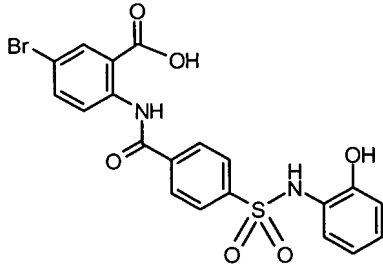
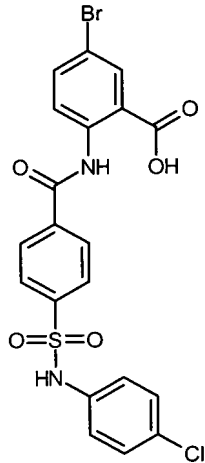
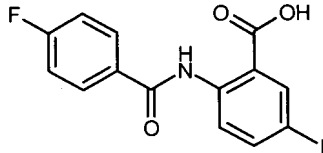
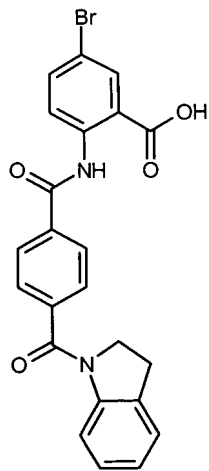
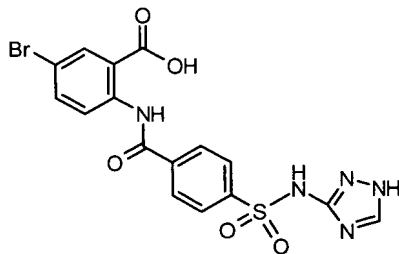
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-656882 	16	PHA-656893 	8
PHA-656884 	16	PHA-662253 	128
PHA-656886 	16	PHA-662412 	64
PHA-656888 	16	PHA-679759 	>128

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-656890 	16	PHA-708922 	>128
PHA-656892 	8	PHA-708977 	>128
PHA-656894 	16	PHA-708987 	>128

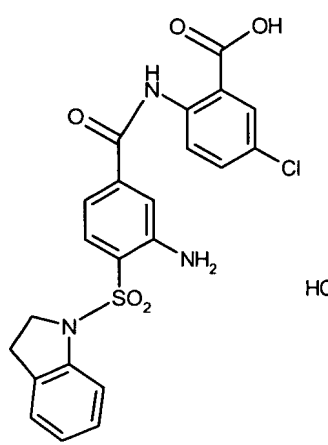
Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
<p>PHA-662254</p> 	>128	<p>PHA-713390</p> 	>128
<p>PHA-679756</p> 	>128	<p>PHA-713392</p> 	>128
<p>PHA-687570</p> 	128	<p>PHA-713395</p> 	>128

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-708979 	>128	PHA-738531 	64
PHA-713389 	>128	PHA-740499 	128
PHA-713391 	>128	PNU-276556 	

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-713393 	>128	PNU-276873 	
PHA-713397 	>128	PNU-282858 	
PHA-738532 	32	PNU-282860 	

Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
PHA-748361 	8	PNU-291997 	1
PNU-276672 		PNU-281164 	>128
PNU-292577 	128	PNU-282859 	32



Compound No., Structure	SA 9218 MIC	Compound No., Structure	SA 9218 MIC
		PNU-290881A  <chem>Nc1cc(cc(c1S(=O)(=O)N2CCc3ccccc32)C(=O)Nc4cc(cc(c4C(=O)O)Cl)C(=O)O)Cl</chem>	4